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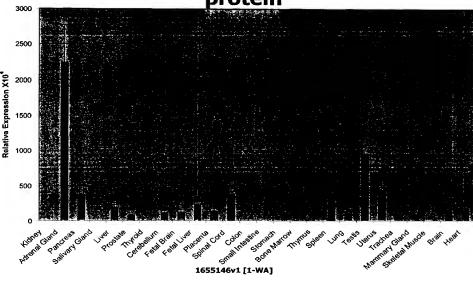
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(54) Title: IDENTIFYING DRUGS FOR AND DIAGNOSIS OF BENIGN PROSTATIC HYPERPLASIA USING GENE EXPRESSION PROFILES

# N91971, cellular retinol binding protein



(57) Abstract: The present invention is based on the elucidation of the global changes in gene expression in prostate tissue isolated from patients exhibiting different clinical states of prostate hyperplasia as compared to normal prostate tissue as well as the identification of individual genes that are differentially expressed in diseased prostate tissue.





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# IDENTIFYING DRUGS FOR AND DIAGNOSIS OF BENIGN PROSTATIC HYPERPLASIA USING GENE EXPRESSION PROFILES

#### RELATED APPLICATIONS

This application claims priority of U.S. Provisional Application No. 60/223,323, filed August 7, 2000, and U.S. Application No. 09/873,319, filed June 5, 2001, which are herein incorporated by reference in their entirety.

#### **BACKGROUND OF THE INVENTION**

Benign Prostatic Hyperplasia (BPH) is the most common benign tumor in men aged >60 years. It is estimated that one in four men living to the age of 80 will require treatment for this disease. BPH is usually noted clinically after the age of 50, the incidence increasing with age, but as many as two thirds of men between the ages of 40 and 49 demonstrate histological evidence of the disease.

The anatomic location of the prostate at the bladder neck enveloping the urethra plays an important role in the pathology of BPH, including bladder outlet obstruction. Two prostate components are thought to play a role in bladder outlet obstruction. The first is the relative increased prostate tissue mass. The second component is the prostatic smooth muscle tone.

The causative factors of BPH in man have been intensively studied. See Ziada *et al.*, *Urology*, 53: 1-6, 1999. In general, the two most important factors appear to be aging and the presence of functional testes. Although these factors appear to be key to the development of BPH, both appear to be nonspecific.

Little is known about the molecular changes in prostate cells associated with the development and progression of BPH. It has been demonstrated that the expression levels of a number of individual genes are changed compared to normal prostate cells. These changes in gene expression include decreased expression of Wilm's tumor gene (WT-1) and increased expression of insulin growth factor II (IGF-II) (Dong *et al.*, *J. Clin. Endocrin. Metab.*, 82(7): 2198-220).

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While the changes in the expression levels of a number of individual genes have been identified, the investigation of the global changes in gene expression has not been reported. Accordingly, there exists a need for the investigation of the changes in global gene expression levels as well as the need for the identification of new molecular markers associated with the development and progression of BPH. Furthermore, if intervention is expected to be successful in halting or slowing down BPH, means of accurately assessing the early manifestations of BPH need to be established. One way to accurately assess the early manifestations of BPH is to identify markers which are uniquely associated with disease progression. Likewise, the development of therapeutics to prevent or stop the progression of BPH relies on the identification of genes responsible for BPH growth and function.

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#### SUMMARY OF THE INVENTION

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The present invention is based on the elucidation of the global changes in gene expression in BPH tissue isolated from patients exhibiting different clinical states of prostate hyperplasia as compared to normal prostate tissue as well as the identification of individual genes that are differentially expressed in BPH tissue.

The invention is also based on the discovery of a means of effectively selecting disease-linked drug targets from gene expression results. The invention includes methods of classifying genes whose expression levels are changed in diseased tissues, during disease induction or during disease progression into specific groups. By using this method it is possible to classify genes whose expression are regulated by the same mechanism into the same group, and it is possible to identify representative marker genes by selecting typical genes from each cluster.

The invention includes methods of screening for or identifying an agent that modulates the onset or progression of BPH, comprising: preparing a first gene expression profile of BPH cells; exposing the cells to the agent; preparing a second gene expression profile of the agent exposed cells; and comparing the first and second gene expression profiles. In a preferred embodiment of these methods, the gene expression profile comprises the expression levels of one or more or preferably two or more genes in Tables 1-5. In another preferred embodiment of these methods, the cell is a prostate cell from a BPH patient, a cell line in Table 6, or a derivative thereof.

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The invention also includes methods of monitoring a treatment of a patient with BPH, comprising administering a pharmaceutical composition to the patient; preparing a gene expression profile from a prostate cell or tissue sample from the patient; and comparing the patient gene expression profile to a gene expression profile from a normal prostate cell population, a BPH tissue or BPH cells without treatment with the pharmaceutical composition. In preferred embodiments of these methods, the gene expression profile comprises the expression levels of one or more or, preferably two or more genes in Tables 1-5.

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The invention also includes methods of diagnosing benign prostatic hyperplasia (BPH) in a subject comprising the step of detecting the level of expression in a tissue or cell sample from the subject of two or more genes from Tables 1-5 (preferably Tables 3-5, and more preferably Table 5); wherein differential expression of the genes is indicative of BPH progression.

The invention further includes methods of detecting the onset or progression of benign prostatic hyperplasia (BPH) in a patient comprising the step of detecting the level of expression in a tissue or cell sample of two or more genes from Tables 1-5 (preferably Tables 3-5, and more preferably Table 5); wherein differential expression of the genes is indicative of BPH progression.

The invention also includes methods of differentiating benign prostatic hyperplasia (BPH) from prostate cancer in a patient comprising the step of detecting the level of expression in a tissue or cell sample of two or more genes from Tables 1-5 (preferably Tables 3-5, and more preferably Table 5); wherein differential expression of the genes is indicative of BPH rather than prostate cancer.

The invention also includes methods of selecting or identifying cells that can be used for drug screening.

All of these methods may include the step of detecting the expression levels of at least about 2, 3, 4, 5, 6, 7, 8, 9, 10 or more genes in any of Tables 1-5, or preferably Table 5. In a preferred embodiment, expression of all of the genes or nearly all of the genes in Tables 1-5, or preferably Table 5, may be detected.

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The invention further includes sets of at least two or more probes, wherein each of the probes comprises a sequence that specifically hybridizes to a gene in Tables 1-5 as well as solid supports comprising at least two or more of the probes.

The invention also includes computer systems comprising or linked to a database containing information identifying the expression level in BPH tissue or cells of a set of genes comprising at least two genes in Tables 1-5, preferably from Table 5; and a user interface to view the information. The database may further comprise sequence information for the genes as well as information identifying the expression level for the set of genes in normal prostate tissue or cells, and prostate cancer tissue. The database may further contain or be linked to descriptive information from an external database, which information correlates said genes to records in the external database.

The invention further includes methods of using the disclosed computer systems to present information identifying the expression level in a tissue or cell of a set of genes comprising at least one of the genes in Tables 1-5, preferably Table 5, comprising comparing the expression level of at least one gene in Tables 1-5, preferably Table 5, in the tissue or cell to the level of expression of the gene in the database.

Lastly, the invention includes kits comprising probes or solid supports of the invention. In some embodiments, the kits also contain written materials or software concerning gene expression information for the genes of the invention, preferably in electronic format.

#### BRIEF DESCRIPTION OF THE DRAWINGS

- Figure 1. Figure 1 shows the expression of cellular retinol binding protein RNA in various tissues.
- Figure 2 shows the expression of cellular retinol binding protein RNA in various prostate tissues samples. In all of the figures, "Normal", "-Sym", "Cancer" and "+Sym" refer to normal prostate, BPH without symptoms, prostate cancer, and BPH with symptoms, respectively.
- Figure 3. Figure 3 shows the expression of S100 calcium binding protein RNA in various tissues.

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Figure 4. Figure 4 shows the expression of S100 calcium binding protein RNA in various prostate tissue samples.

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Figure 5. Figure 5 shows the expression of human prostate-specific membrane antigen (PSMA) RNA in various tissues.

Figure 6. Figure 6 shows the expression of PSMA RNA in various prostate tissue samples.

#### DETAILED DESCRIPTION

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Many biological functions are accomplished by altering the expression of various genes through transcriptional (e.g. through control of initiation, provision of RNA precursors, RNA processing, etc.) and/or translational control. For example, fundamental biological processes such as cell cycle, cell differentiation and cell death, are often characterized by the variations in the expression levels of groups of genes.

Changes in gene expression also are associated with pathogenesis. For example, the lack of sufficient expression of functional tumor suppressor genes and/or the over expression of oncogene/protooncogenes could lead to tumorgenesis or hyperplastic growth of cells (Marshall, Cell, 64: 313-326 (1991); Weinberg, Science, 254:1138-1146 (1991)). Thus, changes in the expression levels of particular genes (*e.g.* oncogenes or tumor suppressors) serve as signposts for the presence and progression of various diseases.

Monitoring changes in gene expression may also provide certain advantages during drug screening development. Often drugs are screened for the ability to interact with a major target without regard to other effects the drugs have on cells. Often such other effects cause toxicity in the whole animal, which prevent the development and use of the potential drug.

The present inventors have examined tissue from normal prostate, BPH and BPH prostate tissue immediately adjacent to malignant prostate tissue to identify the global changes in gene expression in BPH. These changes in gene expression, also referred to as expression profiles, provide useful markers for diagnostic uses as well as markers that can be used to monitor disease states, disease progression, toxicity, drug efficacy and drug metabolism.

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#### Assay Formats

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The genes identified as being differentially expressed in BPH tissue or BPH cells (Tables 1-5) may be used in a variety of nucleic acid detection assays to detect or quantititate the expression level of a gene or multiple genes in a given sample. For example, traditional Northern blotting, nuclease protection, RT- PCR and differential display methods may be used for detecting gene expression levels. Those methods are useful for some embodiments of the invention, particularly when smaller numbers of genes are assayed. For instance, when fewer than 50 genes are assayed, RT-PCR techniques can be used to prepare high-throughput assays. However, methods and assays of the invention are most efficiently designed with hybridization-based methods for detecting the expression of a large number of genes.

Any hybridization assay format may be used, including solution-based and solid support-based assay formats. Solid supports containing oligonucleotide probes for differentially expressed genes of the invention can be filters, polyvinyl chloride dishes, silicon or glass based beads or chips, etc. Such supports and hybridization methods are widely available, for example, those disclosed by Beattie (WO 95/11755). Any solid surface to which oligonucleotides can be bound, either directly or indirectly, either covalently or non-covalently, can be used.

A preferred solid support is a high density array or DNA chip. These contain a particular oligonucleotide probe in a predetermined location on the array. Each predetermined location may contain more than one molecule of the probe, but each molecule within the predetermined location has an identical sequence. Such predetermined locations are termed features. There may be, for example, from 2, 10, 100, 1000 to 10,000, 100,000 or 400,000 of such features on a single solid support. The solid support, or the area within which the probes are attached may be on the order of about a square centimeter.

Oligonucleotide probe arrays for expression monitoring can be made and used according to any technique known in the art (see for example, Lockhart *et al.*, Nat. Biotechnol. (1996) 14, 1675-1680; McGall *et al.*, *Proc. Nat. Acad. Sci. USA* (1996) 93, 13555-13460). Such probe arrays may contain at least two or more oligonucleotides that are complementary to or hybridize to two or more of the genes described in Tables 1-5. For instance, such arrays may contain oligonucleotides that are complementary or hybridize to at least about 2, 3, 4, 5, 6, 7, 8, 9, 10, 20, 30, 50, 70 or more the genes described herein.

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The genes which are assayed according to the present invention are typically in the form of mRNA or reverse transcribed mRNA. The genes may be cloned or not. The genes may be amplified or not. The cloning itself does not appear to bias the representation of genes within a population. However, it may be preferable to use polyA+RNA as a source, as it can be used with less processing steps.

The sequences and related information of the genes described herein are available in the public databases. Tables 1-5 provide the Accession numbers and name for each of the sequences. Each Accession Number corresponds to a sequence in the attached sequence listing. The sequences and related information of the genes listed in the Tables according to their GenBank identifiers are expressly incorporated herein as of the filing date of this application, as are sequences in the databases related to those herein described, such as fragments, variant sequences, etc. (see: <a href="https://www.ncbi.nlm.nih.gov/">www.ncbi.nlm.nih.gov/</a>).

Probes based on the sequences of the genes described above may be prepared by any commonly available method. Oligonucleotide probes for interrogating the tissue or cell sample are preferably of sufficient length to specifically hybridize only to appropriate, complementary genes or transcripts. Typically the oligonucleotide probes will be at least 10, 12, 14, 16, 18, 20 or 25 nucleotides in length. In some cases longer probes of at least 30, 40, or 50 nucleotides will be desirable.

As used herein, oligonucleotide sequences that are complementary to one or more of the genes described in Tables 1-5 refer to oligonucleotides that are capable of hybridizing under stringent conditions to at least part of the nucleotide sequence of said genes. Such hybridizable oligonucleotides will typically exhibit at least about 75% sequence identity at the nucleotide level to said genes, preferably about 80% or 85% sequence identity or more preferably about 90% or 95% or more sequence identity to said genes.

"Bind(s) substantially" refers to complementary hybridization between a probe nucleic acid and a target nucleic acid and embraces minor mismatches that can be accommodated by reducing the stringency of the hybridization media to achieve the desired detection of the target polynucleotide sequence.

The terms "background" or "background signal intensity" refer to hybridization signals resulting from non-specific binding, or other interactions, between the labeled target nucleic acids and components of the oligonucleotide array (e.g., the oligonucleotide probes, control probes, the array substrate, etc.). Background signals may also be produced by

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intrinsic fluorescence of the array components themselves. A single background signal can be calculated for the entire array, or a different background signal may be calculated for each target nucleic acid. In a preferred embodiment, background is calculated as the average hybridization signal intensity for the lowest 5% to 10% of the probes in the array, or, where a different background signal is calculated for each target gene, for the lowest 5% to 10% of the probes for each gene. Of course, one of skill in the art will appreciate that where the probes to a particular gene hybridize well and thus appear to be specifically binding to a target sequence, they should not be used in a background signal calculation. Alternatively, background may be calculated as the average hybridization signal intensity produced by hybridization to probes that are not complementary to any sequence found in the sample (e.g. probes directed to nucleic acids of the opposite sense or to genes not found in the sample such as bacterial genes where the sample is mammalian nucleic acids). Background can also be calculated as the average signal intensity produced by regions of the array that lack probes.

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The phrase "hybridizing specifically to" refers to the binding, duplexing, or hybridizing of a molecule substantially to or only to a particular nucleotide sequence or sequences under stringent conditions when that sequence is present in a complex mixture (e.g., total cellular DNA or RNA).

Assays and methods of the invention may utilize available formats to simultaneously screen at least about 100, preferably about 1000, more preferably about 10,000 and most preferably about 1,000,000 different nucleic acid hybridizations.

As used herein a "probe" is defined as a nucleic acid molecule, capable of binding to a target nucleic acid of complementary sequence through one or more types of chemical bonds, usually through complementary base pairing, usually through hydrogen bond formation. As used herein, a probe may include natural (*i.e.*, A, G, U, C, or T) or modified bases (7-deazaguanosine, inosine, *etc.*). In addition, the bases in probes may be joined by a linkage other than a phosphodiester bond, so long as it does not interfere with hybridization. Thus, probes may be peptide nucleic acids in which the constituent bases are joined by peptide bonds rather than phosphodiester linkages.

The term "stringent conditions" refers to conditions under which a probe will hybridize to its target subsequence, but with only insubstantial hybridization to other sequences or to other sequences such that the difference may be identified. Stringent conditions are sequence-dependent and will be different in different circumstances. Longer

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sequences hybridize specifically at higher temperatures. Generally, stringent conditions are selected to be about 5oC lower than the thermal melting point (Tm) for the specific sequence at a defined ionic strength and pH.

Typically, stringent conditions will be those in which the salt concentration is at least about 0.01 to 1.0 M Na ion concentration (or other salts) at pH 7.0 to 8.3 and the temperature is at least about 30°C for short probes (e.g., 10 to 50 nucleotide). Stringent conditions may also be achieved with the addition of destabilizing agents such as formamide.

The "percentage of sequence identity" or "sequence identity" is determined by comparing two optimally aligned sequences or subsequences over a comparison window or span, wherein the portion of the polynucleotide sequence in the comparison window may optionally comprise additions or deletions (*i.e.*, gaps) as compared to the reference sequence (which does not comprise additions or deletions) for optimal alignment of the two sequences. The percentage is calculated by determining the number of positions at which the identical submit (*e.g.* nucleic acid base or amino acid residue) occurs in both sequences to yield the number of matched positions, dividing the number of matched positions by the total number of positions in the window of comparison and multiplying the result by 100 to yield the percentage of sequence identity. Percentage sequence identity when calculated using the programs GAP or BESTFIT (see below) is calculated using default gap weights.

#### Probe design

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One of skill in the art will appreciate that an enormous number of array designs are suitable for the practice of this invention. The high density array will typically include a number of probes that specifically hybridize to the sequences of interest. See WO 99/32660 for methods of producing probes for a given gene or genes. In addition, in a preferred embodiment, the array will include one or more control probes.

High density array chips of the invention include "test probes." Test probes could be oligonucleotides that range from about 5 to about 500 or 5 to about 45 nucleotides, more preferably from about 10 to about 40 nucleotides and most preferably from about 15 to about 40 nucleotides in length. In other particularly preferred embodiments the probes are 20 or 25 nucleotides in length. In another preferred embodiment, test probes are double or single strand DNA sequences. DNA sequences are isolated or cloned from natural sources or

amplified from natural sources using native nucleic acid as templates. These probes have sequences complementary to particular subsequences of the genes whose expression they are designed to detect. Thus, the test probes are capable of specifically hybridizing to the target nucleic acid they are to detect (the genes of Tables 1-5).

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The term "perfect match probe" refers to a probe that has a sequence that is perfectly complementary to a particular target sequence. The probe is typically perfectly complementary to a portion (subsequence) of the target sequence. The perfect match (PM) probe can be a "test probe", a "normalization control" probe, an expression level control probe and the like. A perfect match control or perfect match probe is, however, distinguished from a "mismatch control" or "mismatch probe."

In addition to test probes that bind the target nucleic acid(s) of interest, the high density array can contain a number of control probes. The control probes fall into three categories referred to herein as 1) normalization controls; 2) expression level controls; and 3) mismatch controls.

Normalization controls are oligonucleotide or other nucleic acid probes that are complementary to labeled reference oligonucleotides or other nucleic acid sequences that are added to the nucleic acid sample to be screened. The signals obtained from the normalization controls after hybridization provide a control for variations in hybridization conditions, label intensity, "reading" efficiency and other factors that may cause the signal of a perfect hybridization to vary between arrays. In a preferred embodiment, signals (e.g., fluorescence intensity) read from all other probes in the array are divided by the signal (e.g., fluorescence intensity) from the control probes thereby normalizing the measurements.

Virtually any probe may serve as a normalization control. However, it is recognized that hybridization efficiency varies with base composition and probe length. Preferred normalization probes are selected to reflect the average length of the other probes present in the array, however, they can be selected to cover a range of lengths. The normalization control(s) can also be selected to reflect the (average) base composition of the other probes in the array, however in a preferred embodiment, only one or a few probes are used and they are selected such that they hybridize well (*i.e.*, no secondary structure) and do not match any target-specific probes.

Expression level controls are probes that hybridize specifically with constitutively expressed genes in the biological sample. Virtually any constitutively expressed gene

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provides a suitable target for expression level controls. Typically expression level control probes have sequences complementary to subsequences of constitutively expressed "housekeeping genes" including, but not limited to an actin gene, the transferrin receptor gene, the GAPDH gene, and the like.

Mismatch controls or mismatch probes may also be provided for the probes to the target genes, for expression level controls or for normalization controls. Mismatch controls are oligonucleotide probes or other nucleic acid probes identical to their corresponding test or control probes except for the presence of one or more mismatched bases. A mismatched base is a base selected so that it is not complementary to the corresponding base in the target sequence to which the probe would otherwise specifically hybridize. One or more mismatches are selected such that under appropriate hybridization conditions (e.g., stringent conditions) the test or control probe would be expected to hybridize with its target sequence, but the mismatch probe would not hybridize (or would hybridize to a significantly lesser extent). Preferred mismatch probes contain a central mismatch. Thus, for example, where a probe is a 20 mer, a corresponding mismatch probe will have the identical sequence except for a single base mismatch (e.g., substituting a G, a C or a T for an A) at any of positions 6 through 14 (the central mismatch).

Mismatch probes thus provide a control for non-specific binding or cross hybridization to a nucleic acid in the sample other than the target to which the probe is directed. Mismatch probes also indicate whether a hybridization is specific or not. For example, if the target is present the perfect match probes should be consistently brighter than the mismatch probes. In addition, if all central mismatches are present, the mismatch probes can be used to detect a mutation. The difference in intensity between the perfect match and the mismatch probe provides a good measure of the concentration of the hybridized material.

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#### Nucleic Acid Samples

As is apparent to one of ordinary skill in the art, nucleic acid samples used in the methods and assays of the invention may be prepared by any available method or process. Methods of isolating total mRNA are well known to those of skill in the art. For example, methods of isolation and purification of nucleic acids are described in detail in Chapter 3 of Laboratory Techniques in Biochemistry and Molecular Biology: Hybridization With Nucleic Acid Probes, Part I Theory and Nucleic Acid Preparation, P. Tijssen, Ed., Elsevier, N.Y.

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(1993). Such samples include RNA samples, but also include cDNA synthesized from a mRNA sample isolated from a cell or tissue of interest. Such samples also include DNA amplified from the cDNA, and RNA transcribed from the amplified DNA. One of skill in the art would appreciate that it is desirable to inhibit or destroy RNase present in homogenates before homogenates can be used.

Biological samples may be of any biological tissue or fluid or cells from any organism as well as cells raised in vitro, such as cell lines and tissue culture cells. Biological samples may also include sections of tissues, such as frozen sections or formalin fixed sections taken for histological purposes. Frequently, the sample will be a "clinical sample" which is a sample derived from a patient. Typical clinical samples include, but are not limited to prostate tissue, urine, sputum, blood, blood-cells (e.g., white cells or peripheral blood leukocytes (PBL), tissue or fine needle biopsy samples, peritoneal fluid, and pleural fluid, or cells therefrom.

#### Forming High Density Arrays.

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Methods of forming high density arrays of oligonucleotides with a minimal number of synthetic steps are known. The oligonucleotide analogue array can be synthesized on a solid substrate by a variety of methods, including, but not limited to, light-directed chemical coupling, and mechanically directed coupling. See Pirrung *et al.*, U.S. Patent No. 5,143, 854.

In brief, the light-directed combinatorial synthesis of oligonucleotide arrays on a glass surface proceeds using automated phosphoramidite chemistry and chip masking techniques. In one specific implementation, a glass surface is derivatized with a silane reagent containing a functional group, e.g., a hydroxyl or amine group blocked by a photolabile protecting group. Photolysis through a photolithogaphic mask is used selectively to expose functional groups which are then ready to react with incoming 5' photoprotected nucleoside phosphoramidites. The phosphoramidites react only with those sites which are illuminated (and thus exposed by removal of the photolabile blocking group). Thus, the phosphoramidites only add to those areas selectively exposed from the preceding step. These steps are repeated until the desired array of sequences have been synthesized on the solid surface. Combinatorial synthesis of different oligonucleotide analogues at different locations on the array is determined by the pattern of illumination during synthesis and the order of addition of coupling reagents.

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In addition to the foregoing, additional methods which can be used to generate an array of oligonucleotides on a single substrate are described WO 93/09668. High density nucleic acid arrays can also be fabricated by depositing premade or natural nucleic acids in predetermined positions. Synthesized or natural nucleic acids are deposited on specific locations of a substrate by light directed targeting and oligonucleotide directed targeting. Another embodiment uses a dispenser that moves from region to region to deposit nucleic acids in specific spots.

#### Hybridization

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Nucleic acid hybridization simply involves contacting a probe and target nucleic acid under conditions where the probe and its complementary target can form stable hybrid duplexes through complementary base pairing. See WO 99/32660. The nucleic acids that do not form hybrid duplexes are then washed away leaving the hybridized nucleic acids to be detected, typically through detection of an attached detectable label. It is generally recognized that nucleic acids are denatured by increasing the temperature or decreasing the salt concentration of the buffer containing the nucleic acids. Under low stringency conditions (e.g., low temperature and/or high salt) hybrid duplexes (e.g., DNA:DNA, RNA:RNA, or RNA:DNA) will form even where the annealed sequences are not perfectly complementary.

Thus specificity of hybridization is reduced at lower stringency. Conversely, at higher stringency (e.g., higher temperature or lower salt) successful hybridization tolerates fewer mismatches. One of skill in the art will appreciate that hybridization conditions may be selected to provide any degree of stringency. In a preferred embodiment, hybridization is performed at low stringency in this case in 6X SSPE-T at 37°C (0.005% Triton X-100) to ensure hybridization and then subsequent washes are performed at higher stringency (e.g., I X SSPE-T at 37°C) to eliminate mismatched hybrid duplexes. Successive washes may be performed at increasingly higher stringency (e.g., down to as low as 0.25 X SSPET at 37°C to 50°C) until a desired level of hybridization specificity is obtained. Stringency can also be increased by addition of agents such as formamide. Hybridization specificity may be evaluated by comparison of hybridization to the test probes with hybridization to the various controls that can be present (e.g., expression level control, normalization control, mismatch controls, etc.).

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In general, there is a tradeoff between hybridization specificity (stringency) and signal intensity. Thus, in a preferred embodiment, the wash is performed at the highest stringency that produces consistent results and that provides a signal intensity greater than approximately 10% of the background intensity. Thus, in a preferred embodiment, the hybridized array may be washed at successively higher stringency solutions and read between each wash. Analysis of the data sets thus produced will reveal a wash stringency above which the hybridization pattern is not appreciably altered and which provides adequate signal for the particular oligonucleotide probes of interest.

#### 10 Signal Detection

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The hybridized nucleic acids are typically detected by detecting one or more labels attached to the sample nucleic acids. The labels may be incorporated by any of a number of means well known to those of skill in the art. See WO 99/32660.

#### 15 Databases

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The present invention includes relational databases containing sequence information, for instance for the genes of Tables 1-5, as well as gene expression information in various prostate tissue samples. Databases may also contain information associated with a given sequence or tissue sample such as descriptive information about the gene associated with the sequence information, metabolic pathway information for the gene or descriptive information concerning the clinical status of the tissue sample, or the patient from which the sample was derived. Such information for the patient may include, but is not limited to sex, age, disease status, general health information, surgical or treatment status, PSA levels, as well as information concerning the patient's clinical symptoms. The database may be designed to include different parts, for instance a sequence database and a gene expression database. Methods for the configuration and construction of such databases are widely available, for instance, see U.S. Patent 5,953,727, which is herein incorporated by reference in its entirety.

The databases of the invention may be linked to an outside or external database. In a preferred embodiment, as described in Tables 1-5, the external database is GenBank and the associated databases maintained by the National Center for Biotechnology Information (NCBI).

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Any appropriate computer platform may be used to perform the necessary comparisons between sequence information, gene expression information and any other information in the database or provided as an input. For example, a large number of computer workstations are available from a variety of manufacturers, such has those available from Silicon Graphics. Client/server environments, database servers and networks are also widely available and appropriate platforms for the databases of the invention.

The databases of the invention may be used to produce, among other things, electronic Northerns that allow the user to determine the cell type or tissue in which a given gene is expressed and to allow determination of the abundance or expression level of a given gene in a particular tissue or cell.

The databases of the invention may also be used to present information identifying the expression level in a tissue or cell of a set of genes comprising at least two of the genes in Tables 1-5, comprising the step of comparing the expression level of at least one gene in Tables 1-5 found or detected in the tissue to the level of expression of the gene in the database. Such methods may be used to predict the hyperplastic state of a given tissue by comparing the level of expression of a gene or genes in Tables 1-5 from a sample to the expression levels found in normal prostate cells, BPH cells or tissue and/or malignant or cancerous prostate tissue. Such methods may also be used in the drug or agent screening assays as described below.

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#### Selection of BPH-Associated Genes

BPH associated genes may be identified or selected by any available method, including subtractive hybridization protocols, differential display protocols and high-throughput hybridization formats, including oligonucleotide and cDNA microarray technologies.

Unprocessed or raw expression levels may be normalized, standardized and/or analyzed by any available computational method, including the expression level normalization, analysis and clustering methods herein described. The normalization method as described in Example 4 may be combined with any further analysis method, including any clustering methods available in the art.

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Diagnostic Uses for the BPH Markers

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As described above, the genes and gene expression information provided in Tables 1-5 may be used as diagnostic markers for the prediction or identification of the hyperplastic state of a prostate or other tissue. For instance, a prostate tissue or other patient sample may be assayed by any of the methods described above, and the expression levels from a gene or genes from Tables 1-5 may be compared to the expression levels found in normal prostate tissue, BPH tissue or BPH tissue from a patient with metastatic or nonmetastatic prostate cancer. In some instances, patient PBLs may be used as the patient sample. The comparison of expression data, as well as available sequence or other information may be done by researcher or diagnostician or may be done with the aid of a computer and databases as described above.

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#### Use of the BPH Markers for Monitoring Disease Progression

As described above, the genes and gene expression information provided in Tables 1-5 may also be used as markers for the monitoring of disease progression, such as the development of BPH. For instance, a prostate tissue or other patient sample may be assayed by any of the methods described above, and the expression levels from a gene or genes from Tables 1-5 may be compared to the expression levels found in normal prostate tissue, BPH tissue or BPH tissue from a patient with metastatic or nonmetastatic prostate cancer. The comparison of the expression data, as well as available sequence or other information may be done by researcher or diagnostician or may be done with the aid of a computer and databases as described above.

The BPH markers of the invention may also be used to track or predict the progress or efficacy of a treatment regime in a patient. For instance, a patient's progress or response to a given drug may be monitored by creating a gene expression profile from a tissue or cell sample after treatment or administration of the drug. The gene expression profile may then be compared to a gene expression profile prepared from normal cells or tissue, for instance, normal prostate tissue. The gene expression profile may also be compared to a gene expression profile prepared from BPH or malignant prostate cells, or from tissue or cells from the same patient before treatment. The gene expression profile may be made from at least one gene, preferably more than one gene, and most preferably all or nearly all of the genes in Tables 1-5.

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#### Use of the BPH Markers for Drug Screening

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According to the present invention, the genes identified in Tables 1-5 can be used as markers to screen for potential therapeutic agents or compounds to treat BPH or prostate cancer. A candidate drug or agent can be screened for the ability to stimulate the transcription or expression of a given marker or to down-regulate or counteract the transcription or expression of a marker or markers. Compounds that modulate the expression level of single gene and also compounds that modulate the expression level of multiple genes from levels associated with a specific disease state to a normal state can be screened by using the markers and profiles identified herein.

According to the present invention, one can also compare the specificity of drug's effects by looking at the number of markers which are differentially expressed after drug exposure and comparing them. More specific drugs will have less transcriptional targets. Similar sets of markers identified for two drugs may indicate a similarity of effects.

Assays to monitor the expression of a marker or markers as defined in Tables 1-5 may utilize any available means of monitoring for changes in the expression level of the nucleic acids of the invention. As used herein, an agent is said to modulate the expression of a nucleic acid of the invention if it is capable of up- or down-regulating expression of the nucleic acid in a cell.

In one assay format, gene chips containing probes to at least 2 genes from Tables 1-5 may be used to directly monitor or detect changes in gene expression in the treated or exposed cell as described in more detail above. In another format, the changes of mRNA expression level can be detected using QuantiGene technology (Warrior *et. al.* (2000) *J. Biomolecular Screening*, 5, 343-351). Specific probes used for QuantiGene can be designed and synthesized to one or more genes from Tables 1-5. Cells treated with compounds are lysed by lysis buffer. The amount of target mRNA can be detected as a luminescence intensity using target specific probes.

In another format, cell lines that contain reporter gene fusions between the open reading frame and/or 5'/3' regulatory regions of a gene in Tables 1-5 and any assayable fusion partner may be prepared. Numerous assayable fusion partners are known and readily available including the firefly luciferase gene and the gene encoding chloramphenicol

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acetyltransferase (Alam et al. (1990) Anal. Biochem. 188:245-254). Cell lines containing the reporter gene fusions are then exposed to the agent to be tested under appropriate conditions and time. Differential expression of the reporter gene between samples exposed to the agent and control samples identifies agents which modulate the expression of the nucleic acid.

Additional assay formats may be used to monitor the ability of the agent to modulate the expression of a gene identified in Tables 1-5. For instance, as described above, mRNA expression may be monitored directly by hybridization of probes to the nucleic acids of the invention. Cell lines are exposed to the agent to be tested under appropriate conditions and time and total RNA or mRNA is isolated by standard procedures such those disclosed in Sambrook *et al.* (*Molecular Cloning: A Laboratory Manual*, 2nd Ed. Cold Spring Harbor Laboratory Press, 1989).

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In another assay format, cells or cell lines are first identified which express the gene products of the invention physiologically (see below). Cell and/or cell lines so identified would be expected to comprise the necessary cellular machinery such that the fidelity of modulation of the transcriptional apparatus is maintained with regard to exogenous contact of agent with appropriate surface transduction mechanisms and/or the cytosolic cascades. Such cell lines may be, but are not required to be, prostate derived. Further, such cells or cell lines may be transduced or transfected with an expression vehicle (e.g., a plasmid or viral vector) construct comprising an operable non-translated 5'-promoter containing end of the structural gene encoding the instant gene products fused to one or more antigenic fragments, which are peculiar to the instant gene products, wherein said fragments are under the transcriptional control of said promoter and are expressed as polypeptides whose molecular weight can be distinguished from the naturally occurring polypeptides or may further comprise an immunologically distinct tag or some other detectable marker or tag. Such a process is well known in the art (see Maniatis).

Cells or cell lines transduced or transfected as outlined above are then contacted with agents under appropriate conditions; for example, the agent comprises a pharmaceutically acceptable excipient and is contacted with cells comprised in an aqueous physiological buffer such as phosphate buffered saline (PBS) at physiological pH, Eagles balanced salt solution (BSS) at physiological pH, PBS or BSS comprising serum or conditioned media comprising PBS or BSS and/or serum incubated at 37°C. Said conditions may be modulated as deemed necessary by one of skill in the art. Subsequent to contacting the cells with the agent, said

cells are disrupted and the polypeptides of the lysate are fractionated such that a polypeptide fraction is pooled and contacted with an antibody to be further processed by immunological assay (e.g., ELISA, immunoprecipitation or Western blot). The pool of proteins isolated from the "agent-contacted" sample is then compared with a control sample where only the excipient is contacted with the cells and an increase or decrease in the immunologically generated signal from the "agent-contacted" sample compared to the control is used to distinguish the effectiveness of the agent.

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Another embodiment of the present invention provides methods for identifying agents that modulate at least one activity of a protein(s) encoded by the genes in Tables 1-5. Such methods or assays may utilize any means of monitoring or detecting the desired activity.

In one format, the relative amounts of a protein of the invention between a cell population that has been exposed to the agent to be tested compared to an un-exposed control cell population may be assayed. In this format, probes such as specific antibodies are used to monitor the differential expression of the protein in the different cell populations. Cell lines or populations are exposed to the agent to be tested under appropriate conditions and time. Cellular lysates may be prepared from the exposed cell line or population and a control, unexposed cell line or population. The cellular lysates are then analyzed with the probe, such as a specific antibody.

Agents that are assayed in the above methods can be randomly selected or rationally selected or designed. As used herein, an agent is said to be randomly selected when the agent is chosen randomly without considering the specific sequences involved in the association of the a protein of the invention alone or with its associated substrates, binding partners, *etc*. An example of randomly selected agents is the use a chemical library or a peptide combinatorial library, or a growth broth of an organism.

As used herein, an agent is said to be rationally selected or designed when the agent is chosen on a nonrandom basis which takes into account the sequence of the target site and/or its conformation in connection with the agent's action. Agents can be rationally selected or rationally designed by utilizing the peptide sequences that make up these sites. For example, a rationally selected peptide agent can be a peptide whose amino acid sequence is identical to or a derivative of any functional consensus site.

The agents of the present invention can be, as examples, peptides, small molecules, vitamin derivatives, as well as carbohydrates. Dominant negative proteins, DNAs encoding

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these proteins, antibodies to these proteins, peptide fragments of these proteins or mimics of these proteins may be introduced into cells to affect function. "Mimic" used herein refers to the modification of a region or several regions of a peptide molecule to provide a structure chemically different from the parent peptide but topographically and functionally similar to the parent peptide (see Grant GA. in: Meyers (ed.) Molecular Biology and Biotechnology (New York, VCH Publishers, 1995), pp. 659-664). A skilled artisan can readily recognize that there is no limit as to the structural nature of the agents of the present invention.

#### Cells used for Multi Gene Screening

Many kinds of cells such as primary cells and cell lines can be used for the drug screening methods of the invention. Cells or cell lines derived from prostatic tissues are preferred because the innate gene expression mechanisms of these cells often resemble those of prostatic tissues. Cells used for drug screening can be selected by assaying for the expression of one or more of the marker genes listed in Tables 1-5. The cells which differentially express one or more, or preferably nearly all of the marker genes listed in Tables 1-5 are preferred cells or cell lines for the methods of the invention (see Table 6).

Kits

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The invention further includes kits combining, in different combinations, high-density oligonucleotide arrays, reagents for use with the arrays, signal detection and array-processing instruments, gene expression databases and analysis and database management software described above. The kits may be used, for example, to diagnose the disease state of a tissue or cell sample, to monitor the progression of prostate disease states, to identify genes that show promise as new drug targets and to screen known and newly designed drugs as discussed above.

The databases packaged with the kits are a compilation of expression patterns from human and laboratory animal genes and gene fragments (corresponding to the genes of Tables 1-5). In particular, the database software and packaged information include the expression results of Tables 1-5 that can be used in the assays and methods as herein described. In another format, database access is provided to the purchaser or user through an electronic means, *e.g.*, via the Internet or by direct dial-in access.

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The kits may used in the pharmaceutical industry, where the need for early drug testing is strong due to the high costs associated with drug development, but where bioinformatics, in particular gene expression informatics, is still lacking. These kits will reduce the costs, time and risks associated with traditional new drug screening using cell cultures and laboratory animals. The results of large-scale drug screening of pre-grouped patient populations, pharmacogenomics testing, can also be applied to select drugs with greater efficacy and fewer side-effects. The kits may also be used by smaller biotechnology companies and research institutes who do not have the facilities for performing such large-scale testing themselves.

Databases and software designed for use with use with microarrays is discussed in Balaban *et al.*, U.S. Patent Nos. 6,229,911, a computer-implemented method for managing information, stored as indexed tables, collected from small or large numbers of microarrays, and 6,185,561, a computer-based method with data mining capability for collecting gene expression level data, adding additional attributes and reformatting the data to produce answers to various queries. Chee *et al.*, U.S. Patent No. 5,974,164, disclose a software-based method for identifying mutations in a nucleic acid sequence based on differences in probe fluorescence intensities between wild type and mutant sequences that hybridize to reference sequences.

Without further description, it is believed that one of ordinary skill in the art can, using the preceding description and the following illustrative examples, make and utilize the genes, chips, *etc.* of the present invention and practice the claimed methods. The following working examples therefore, specifically point out the preferred embodiments of the present invention, and are not to be construed as limiting in any way the remainder of the disclosure.

#### EXAMPLES

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#### Example 1: Gene chip expression analysis

Human tissue was obtained from the transitional zone of the prostate (the junction between the ejaculatory duct and the prostatic urethra) in biopsy samples from normal individuals and from patients with BPH or prostate cancer. BPH was defined histologically in all samples. Normal tissue and asymptomatic BPH samples came from individuals who died of trauma and did not report symptoms. Because BPH is a disease associated with aging, two groups of normal individuals were identified, group 1, ages 20 or under, and group

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2, ages 30-50. Patients having BPH with symptoms were defined as those with a need for frequent urination. In these patients, a radical prostatectomy had been performed. Prostate cancer patients provided age-matched tissue samples for symptomatic BPH patients, but were without symptoms and without cancer in the transitional zone under histological examination.

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Microarray sample preparation was conducted with minor modifications, following the protocols set forth in the Affymetrix GeneChip Expression Analysis Manual. Frozen tissue was ground to a powder using a Spex Certiprep 6800 Freezer Mill. Total RNA was extracted with Trizol (GibcoBRL) utilizing the manufacturer's protocol. The total RNA yield for each sample was 200-500 μg per 300 mg tissue weight. mRNA was isolated using the Oligotex mRNA Midi kit (Qiagen) followed by ethanol precipitation. Double stranded cDNA was generated from mRNA using the SuperScript Choice system (GibcoBRL). First strand cDNA synthesis was primed with a T7-(dT24) oligonucleotide. The cDNA was phenol-chloroform extracted and ethanol precipitated to a final concentration of 1 μg/ml. From 2 μg of cDNA, cRNA was synthesized using Ambion's T7 MegaScript *in vitro* Transcription Kit.

To biotin label the cRNA, nucleotides Bio-11-CTP and Bio-16-UTP (Enzo Diagnostics) were added to the reaction. Following a 37°C incubation for six hours, impurities were removed from the labeled cRNA following the RNeasy Mini kit protocol (Qiagen). cRNA was fragmented (fragmentation buffer consisting of 200 mM Tris-acetate, pH 8.1, 500 mM KOAc, 150 mM MgOAc) for thirty-five minutes at 94°C. Following the Affymetrix protocol, 55 μg of fragmented cRNA was hybridized on the Affymetrix Human 42K array set for twenty-four hours at 60 rpm in a 45°C hybridization oven. The chips were washed and stained with Streptavidin Phycoerythrin (SAPE) (Molecular Probes) in Affymetrix fluidics stations. To amplify staining, SAPE solution was added twice with an anti-streptavidin biotinylated antibody (Vector Laboratories) staining step in between. Hybridization to the probe arrays was detected by fluorometric scanning (Hewlett Packard Gene Array Scanner). Data was analyzed using Affymetrix GeneChip version 3.0 and Expression Data Mining Tool (EDMT) software (version 1.0).

Differential expression of genes between the BPH and normal prostate samples were determined using the Affymetrix GeneChip human gene chip set by the following criteria: 1) For each gene, Affymetrix GeneChip average difference values were determined by standard Affymetrix EDMT software algorithms, which also made "Absent" (=not specifically

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detected as gene expression), "Present" (=detected) or "Marginal" (=not clearly Absent or Present) calls for each GeneChip element; 2) all AveDiff values which were less than +20 (positive 20) were raised to a floor of +20 so that fold change calculations could be made where values were not already greater than or equal to +20; 3) median levels of expression were compared between the normal control group and the BPH with symptoms disease group to obtain greater than or equal 2-fold up/down values; 4) The median value for the higher expressing group needed to be greater or equal to 200 average difference units in order to be considered for statistical significance; 5) Genes passing the criteria of #1-4 were analyzed for statistical significance using a two-tailed T test and deemed statistically significant if p < 0.05. Tables 1 and 2 list the genes and their levels of differential expression (compared to normal samples) in BPH tissue from patients with symptoms of BPH and in BPH tissue immediately adjacent to malignant prostate tissue isolated from male patients.

#### Example 2: Expression profile analysis

Gene expression profiles between normal sample and BPH patient samples were determined by using the following samples: 10 normal; 7 BPH without symptoms; 8 BPH with cancer; and 8 BPH with symptoms. Gene expression profiles were prepared using the 42K Affymetrix Gene Chip set. The methods used were the same as described in Example 1 with the exception of the criteria to select the marker genes.

The criteria used in this study were as follows; 1) For each gene, Affymetrix
GeneChip average difference values were determined by standard Affymetrix EDMT
software algorithms, which also made "Absent" (=not specifically detected as gene
expression), "Present" (=detected) or "Marginal" (=not clearly Absent or Present) calls for
each GeneChip element; 2) all AveDiff values which were less than +20 (positive 20) were
raised to a floor of +20 so that fold change calculations could be made where values were not
already greater than or equal to +20; 3) mean levels of expression were compared between
the normal control group and the BPH with symptoms disease group; 4) genes were arranged
by the fold change starting with the largest one (Fold change calculation was determined by
using logarithmic values in Example 2); and 5) the top 200 up-regulated genes and bottom
200 down-regulated genes were selected. The genes identified in this study are listed in
Tables 3 (normal vs. BPH with symptoms, up regulated) and 4 (normal vs. BPH with
symptoms, down regulated, values are negative fold-change from normal).

Example 3: Selection of Cell lines used for Multi Gene Screening

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A number of cultured cell lines were tested to determine the similarity in gene expression profiles to BPH tissue. Cells were cultured in 6-well plates using the appropriate medium for each cell line. After reaching 90% confluency, cells were lysed with Trizol (GiboBRL) and total RNA was extracted. mRNA was then isolated, cDNA and cRNA was synthesized, and gene expression levels were determined by the Affymetrix Human 42K. Gene Chip set as described in more detail above.

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The gene expression profiles were compared with those of prostatic tissue samples. A panel of 61 genes whose expression levels were up-regulated in BPH with symptoms compared with normal samples and with small variation among samples (within BPH samples and within normal samples) were assayed. The group of genes whose signal intensity was more than 100 in each cell line is summarized in Table 1. A panel of 43 genes whose expression levels were down-regulated in BPH patient with small variation among samples was also assayed. The group of genes whose signal intensity in Affymetrix Gene Chip was "Present call" is also included in Table 1. Similarly, genes whose expression level is up- or down-regulated in patients with BPH and cancer, compared to normal controls, are listed in Table 2.

Forty-eight to 58% of genes applied for this analysis were expressed in the cell lines of Table 6. These results indicate that cell lines, BRF-55T (Biological Research Faculty & Facility Inc.), PZ-HPV7 (ATCC; CRL-2221), BPH-1 (S.W. Hayward *et al.*, *In Vitro Cell Dev. Biol.* 31A, 14-24, 1995) and LNCaP (ATCC; CRL-1740) can be used as a BPH – like cell population to screen for compounds which are capable of modulating gene expression profiles from the disease state to a normal state using the genes of Tables 1-5. In particular, BRF-55T is a useful cell line for screening in the assays of the invention, because 58% genes of the assayed genes were differentially expressed in BRF-55T as compared to BPH with symptoms tissue.

#### Example 4: Cluster analysis of up- or down-regulated genes in BPH

Cluster analysis of the expression results from a large number of genes is often problematic due to variations in the standardization of the gene expression data. To compensate for these variations, a subset of differentially expressed genes was selected by a modified analysis procedure.

In a first step, a gene list comparing normal vs. disease samples was generated by two kinds of comparisons. First, genes were selected that displayed a greater than or equal to

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mean 2-fold up or down regulation using average difference expression values and with p<0.05. Second, genes were selected by ANOVA comparing the normal group of samples with the disease group and with a t value of >3 in the up or down direction. These lists were then combined to create an expression profile characteristic of normal controls and one characteristic of disease in which specific genes are found to be up or down regulated in disease when compared with normal controls.

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In preparation for clustering analysis to identify subgroups of genes that show statistically similar expression patterns, average difference values for the selected genes were normalized across all samples (normal and disease combined) using the following formula:

 $Normalization\ data = (X - Xmean)/Sx$ 

Where Sx is variance (:STD)

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This converts the mean expression value for each gene to 0 and the high and low values to 1 and -1, respectively. Thus, genes with high absolute expression values when compared with genes with low absolute expression values would not skew the comparisons when clustering algorithms are applied.

The measurement of the cluster space distance was determined by using the correlation coefficient (1-r) method and clustering was performed using Ward's method (Ward,J.H. (1963) *Journal of American Statistical Association*, 58. 236.)

The clustering was validated by observing whether multiple elements representing the same genes showing the same direction of expression change (*i.e.*, either up or down) tend to cluster together. To test this standardization and clustering protocol, the expression levels for genes that are represented by more than one element on the 42K gene chip set were analyzed to determine whether the multiple elements for a single gene could be clustered together. For example, tryptase, also known as alpha tryptase or beta (tryptase II) is represented by two separate elements on the 42K human gene chip. This gene is registered with 2 different element names 41268 (5), M33493\_s\_at (code name, Up-170) and 26389 (3), rc AA131322 s at (code name, Up-010).

It was found that the best analysis means for decreasing measurement errors between these two elements is by the Ward method as it gave the most consistent results when compared to other clustering methods. These analysis methods may be incorporated into software or computer readable storage media for storing a computer programmer software.

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#### Example 5: Selection of 60 Marker Genes

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A panel of 60 representative marker genes (listed in Table 5) out of 400 marker genes listed in Tables 3 and 4 can be used in the assays and methods of the invention. The 60 marker genes were selected based on following criteria: (1) expression level is changed greatly in BPH patient samples compared with normal samples; (2) variation of expression levels within BPH samples and within normal samples is small; and (3) expression levels resembling BPH with symptoms are detected in cell line BRF-55T.

#### Example 6: Gene Expression Analysis of Select Genes

The expression levels of three genes from Tables 1-5 (the genes encoding cellular retinol binding protein, S100 calcium binding protein and PSMA) were assayed in various tissues and prostate samples by PCR as described in Example 7 (see Figures 1-6). Each sample was assayed for the level of GAPDH and mRNA corresponding to cellular retinol binding protein, S100 calcium binding protein or PSMA. As seen in Figures 1-6, these three genes are differentially regulated or expressed in BPH tissue from patients with or without symptoms and from BPH tissue from patients with prostate cancer (compared to normal prostate tissue). All three genes are therefore useful markers in the assays of the invention, such as the assays to measure the effect of an agent on BPH or the assays to detect or diagnose the occurrence or progression of BPH.

#### Example 7: Drug Screening Assays

The expression profiles for normal controls and disease samples described above can be used to identify compound hits from a compound library. A hit may be, but is not necessarily, defined as one of three kinds of results:

- 1) The expression of an individual gene is changed in the direction of normal (i.e., if up in disease, then down=hit, if down in disease, then up=hit). The stronger the modulation of an individual gene to a normal phenotype, the stronger the hit status for the compound against that gene.
- 2) The expression of genes that subcluster together is evaluated for an overall pattern of modulation to a normal expression profile. The more genes in a subcluster that are modulated to a normal phenotype, the stronger the hit status for the compound against that subcluster. A subcluster may represent common or interacting cellular pathways.

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3) The overall expression profile of all of the genes being screened is evaluated for modulation to normal. The more genes that are modulated to a normal phenotype, the stronger the hit status for the compound against the entire gene set.

As described above, if a compound modulates the gene expression pattern of the screening system cells more towards any disease phenotype, then it can be used as a molecular probe to find binding proteins and/or define disease-associated cellular pathways.

As an example, candidate agents and compounds are screened for their ability to modulate the expression levels of cellular retinol binding protein, \$100 calcium binding protein and PSMA by exposing a prostate cell line or cell line from BPH tissue to the agent and assaying the expression levels of these genes by real time PCR. Real time PCR detection is accomplished by the use of the ABI PRISM 7700 Sequence Detection System. The 7700 measures the fluorescence intensity of the sample each cycle and is able to detect the presence of specific amplicons within the PCR reaction. Each sample is assayed for the level of GAPDH and mRNA corresponding to cellular retinol binding protein, S100 calcium binding protein and PSMA. GAPDH detection is performed using Perkin Elmer part#402869 according to the manufacturer's directions. Primers were designed for the three genes by using Primer Express, a program developed by PE to efficiently find primers and probes for specific sequences ((1) N91971 - FAM PROBE Forward: 5'- CAT ggC TTT gTT TTA AgA AAA ggA A -3'; Reverse: 5'- AgC CAC CCC CAg gCA T -3'; Probe: 5'-FAM - AgT gAC AAA gCC AAg AgA CAg ACT CTg CTA ACA - TAMRA-3'; (2) X65614 - SYBR: Forward: 5'- AAA gAC AAg gAT gCC gTg gAT -3'; Reverse 5'-AgC CAC gAA CAC gAT gAA CTC-3'; (3) M99487-SYB; Forward 5'-Tgg CTC AgC ACC ACC Aga T-3'; Reverse: 5'-TTC Cag TAA AgC Cag gTC CAA-3')

These primers are used in conjunction with SYBR green (Molecular Probes), a nonspecific double stranded DNA dye, to measure the expression level mRNA corresponding to the genes, which is normalized to the GAPDH level in each sample.

Normalized expression levels from cells exposed to the agent are then compared to the normalized expression levels in control cells. Agents that modulate the expression of one or more the genes may be further tested as drug candidates in appropriate BPH *in vitro* or *in vivo* models.

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The expression profiles or one or more of the individual genes of Tables 1-5 are used as molecular or diagnostic markers to evaluate the disease status of a patient sample. In one embodiment, a patient prostate tissue sample is processed as described herein to produce total cellular or mRNA. The RNA is hybridized to a chip continuing probes that specifically hybridize to one or more, or two or more of the genes in Tables 1-5. The overall expression profile generated, or the expression levels of individual genes are then compared to the profiles as described in Tables 1-5 to determine the disease or hyperplastic state of the patient sample.

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Although the present invention has been described in detail with reference to examples above, it is understood that various modifications can be made without departing from the spirit of the invention. Accordingly, the invention is limited only by the following claims. All cited patents, applications, GenBank Accession numbers and publications referred to in this application are herein incorporated by reference in their entirety.

Normal1-Norma	al2 vs BPH-With Symp	toms Table	29 TABLE 1		
1654533.1		Genbank	Genbank	Fold-change	p-value
	Affy element	ID	Name	N1-N2 vs With	N1-N2 vs With
	<b>,</b>		B-cell-homing chemokine (ligand for		
Up-regulated	RC AA410383_at	AA410383	Burkitt's lymphoma receptor-1)4g21	22.5	0.025197485
op rogulatou	RC AA463726_s at	AA463726	JM27 proteinXp11.23	14.9	
	110_747400720_0_0	701400720	Homo sapiens mRNA; cDNA	14.5	0.010000171
			· · · · · · · · · · · · · · · · · · ·		
	RC_AA057195_at	A A 057405	DKFZp586M121 (from clone	14.0	0.000005045
	RC_AA057 195_at	AA057195	DKFZp586M121)	14.0	0.029325045
	1/04540 mad at	1/04540	v-fos FBJ murine osteosarcoma viral	40.4	0.004.007504
	V01512_rna1_at	V01512_rna1	oncogene homolog14q24.3	13.1	0.001027561
	RC_AA427622_s_at	AA427622	collagen, type XIII, alpha 110q22	11.6	0.00074954
			v-fos FBJ murine osteosarcoma viral		
	RC_N23730_s_at	N23730	oncogene homolog14q24.3	11.4	
	RC_AA465491_at	AA465491	Mad4 homolog4p16.3	11.4	0.031024189
	RC_AA620825_at	AA620825	ESTs	11.3	0.010915901
	RC_R93908_at	R93908	ESTs	11.3	0.019994337
	RC_AA461300_at	AA461300	ESTs	11.0	0.007061759
	N40141 at	N40141	JM27 proteinXp11.23	10.9	0.013756347
	RC R25410 at	R25410	ESTs	7.7	0.01851753
			FBJ murine osteosarcoma viral		
	L49169 at	L49169	oncogene homolog B19q13.3	7.4	0.041523744
	RC AA279760 at	AA279760	ESTs	7.0	
	RC_T90889_at	T90889	ESTs	6.5	
	NC_190009_at	190009		0.5	0.015666863
	LICOME -t	LICODAT	insulin-like growth factor binding	0.0	0.000040004
	U62015_at	U62015	protein 101p22-p31	6.0	0.002843661
			highly expressed in cancer, rich in		
	RC_AA188981_at	AA188981	leucine heptad repeats	5.9	
	D83018_at	D83018	nel (chicken)-like 212q13.11-q13.12	5.6	0.000570952
			immunoglobulin gamma 3 (Gm		
	RC_H64493_f_at	H64493	marker)14q32.33	5.6	0.01109802
	X52541_at	X52541	early growth response 15q31.1	5.2	0.002428259
			major histocompatibility complex,		
	M57466 s at	M57466	class II, DP beta 16p21.3	5.1	0.002137399
	J03507 at	J03507	complement component 75p13	4.9	1.36616E-05
	RC_N30198 at	N30198	ESTs	4.8	
	RC T78398 at	T78398	EST	4.8	0.033293747
	RC_H17550_at	H17550	ESTs	4.7	
	110_1111000_dt	1117000		4.7	0.047020022
	RC_T67053_f_at	T67053	immumoglobulin lambda gene	4.5	0.045107075
	RC_AA598982 s at	AA598982	cluster22q11.1-q11.2	4.5	
			trophininXp11.22-p11.21	4.3	
	RC_AA256268_at	AA256268	ESTs	4.2	0.001506239
	1100540115555		insulin-like growth factor 2		
	HG3543-HT3739_at	M29645	(somatomedin A)11p15.5	4.1	0.017253126
	RC_N91971_f_at	N91971	retinol-binding protein 1, cellular3q23	4.1	0.02528773
	RC_AA479286_at	AA479286	ESTs	4.0	0.028009544
	M62831_at	M62831	immediate early protein19	4.0	0.000484086
			ESTs, Weakly similar to unknown		
	RC_F02992_at	F02992	[M.musculus]	3.9	0.031845412
	RC_H86112_f_at	H86112	KIAA0471 gene product1q24-q25	3.8	0.004155259
	RC_AA436616_at	AA436616	ESTs	. 3.8	0.017156387
	RC T62857 at	T62857	ESTs	3.7	0.000301735
	RC_AA281345_f_at	AA281345	immediate early protein19	3.6	0.001679723
	U21128 at	U21128	lumican12q21.3-q22	3.6	2.19529E-05
	U30521 at	U30521	P311 protein	3.6	0.001150397
	RC_N58172_at	N58172	ESTs	3.5	
	RC_T03229_f_at				
	RC_103229_1_at	T03229	EST	3.5	0.031101935
			collagen, type III, alpha 1 (Ehlers-		
	V06700 - at	V06700	Danlos syndrome type IV, autosomal		0.000.45555
	X06700_s_at	X06700	dominant)2q31	3.5	0.008472599
	DO 700004	700004	Homo sapiens clone 23555 mRNA		0.6
	RC_Z39904_at	Z39904	sequence	3.4	0.002949046
	RC_T23622_at	T23622	ESTs	3.4	0.002174281
			immunoglobulin gamma 3 (Gm		
	J00231_f_at	J00231	marker)14q32.33	3.4	0.009322568
	RC_AA028092_s_at	AA028092	transcription factor 216pter-gter	3.4	3.13963E-06
	RC_AA252528_at	AA252528	ESTs	3.4	0.000225707
			procollagen C-endopeptidase		
	L33799_at	L33799	enhancer7q22	3.3	0.018469201
	<del>-</del>		•		•

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	al2 vs BPH-With Symp		TABLE 1		
1654533.		Genbank	Genbank	Fold-change	p-value
	Affy element	ID	Name	N1-N2 vs With	N1-N2 vs With
			Homo sapiens mRNA; cDNA		
			DKFZp586K1220 (from clone		
	RC_F09748_s_at	F09748	DKFZp586K1220)	3.2	0.02728166
			carboxypeptidase A3 (mast cell)3q21		
	RC_T64223_s_at	T64223	q25	3.2	0.027915742
			immunoglobulin gamma 3 (Gm		
	RC_AA402903_f_at	AA402903	marker)14q32.33	3.2	0.044721116
	RC_F13763_at	F13763	ESTs	3.1	0.000503701
	RC_AA488432_at	AA488432	phosphoserine phosphatase7p21-	3.1	0.020997503
			small inducible cytokine A5		
	RC_AA486072_i_at	AA486072	(RANTES)17q11.2-q12	3.1	0.025877597
	RC_N22006_s_at	N22006	ÈST	3.1	0.00148561
	RC_AA257093_r_at	AA257093	T-cell receptor, beta cluster7q35	3.1	1.71945E-07
	RC_AA609943_at	AA609943	ESTs	3.0	0.029360518
	RC_T23490_s_at	T23490	ESTs	3.0	
	D13628_at	D13628	angiopoletin 18q22.3-q23	2.9	0.006228419
			carboxypeptidase A3 (mast cell)3q21		
	M73720 at	M73720	q25	2.9	0.006585391
	Z74616 s at	Z74616	collagen, type I, alpha 27q22.1	2.8	
	AA082546 at	AA082546	ESTs	2.8	
	RC_AA284920 at	AA284920	ESTs	2.7	
	RC AA599365 at	AA599365	decorin12q23	2.7	
	110_11.000000_at	7 11 1000000	insulin-like growth factor 1	2.1	0.001233330
	X57025 at	X57025	3	2.7	0.000044404
	X51345_at	X51345	(somatomedin C)12q22-q23		
	701045_at	A01040	jun B proto-oncogene19p13.2	2.7	0.036487159
	DC NETOTE a at	NETOTE	insulin-like growth factor 1	0.7	0.005040404
	RC_N67876_s_at RC_AA609504 at	N67876	(somatomedin C)12q22-q23	2.7	
	RC_AA009504_at	AA609504	KIAA0405 gene product	2.7	0.020881055
			ESTs, Moderately similar to !!!! ALU		
	DO NG0007 -4	Nonna	SUBFAMILY SB2 WARNING		
	RC_N69207_at	N69207	ENTRY !!!! [H.sapiens]	2.6	0.041315387
			immunoglobulin gamma 3 (Gm		
	M87789_s_at	M87789	marker)14q32.33	2.6	0.038916248
			nuclear receptor subfamily 2, group		
	HG3510-HT3704_at	X12795	F, member 15q14	2.6	0.016151338
			ESTs, Weakly similar to pancortin-1		
	RC_T64211_at	T64211	[M.musculus]	2.6	0.006233291
			butyrophilin, subfamily 3, member		
	U90552_s_at	U90552	A16p23	2.6	0.004564282
			immunoglobulin lambda-like		
	M34516_r_at	M34516	polypeptide 322q11.2	2.6	0.049767038
	RC_T23468_at	T23468	ESTs	2.5	0.00250737
	_		ESTs, Weakly similar to !!!! ALU		
			SUBFAMILY SQ WARNING ENTRY		
	RC AA173223 at	AA173223	!!!! [H.sapiens]	2.5	0.007080285
	RC T49061 at	T49061	ESTs	2.5	0.039642391
	RC_AA234095 at	AA234095	ESTs	2.5	0.003152859
			pre-B-cell leukemia transcription		0.000.02000
	RC_F01920_s_at	F01920	factor 39q33-q34	2.5	0.002088945
	RC N91461 at	N91461	ESTs	2.4	0.01015467
	RC_N67575 s at	N67575	osteoglycin (osteoinductive factor)	2.4	0.004044061
	RC_AA151210_at	AA151210	ESTs	2.4	0.011476541
		701101210	Homo sapiens mRNA; cDNA	2.4	0.011470541
			DKFZp564I1922 (from clone		
	AA156897_s_at	AA156897		2.4	0.00074004
	W73859 at	W73859	DKFZp564I1922)	2.4	0.033974981
	RC_H68097 at	H68097	transcription factor 216pter-qter	2.4	0.024640626
	RC_AA436618 at	AA436618	EST	2.4	0.04870874
	M33493_s at		ESTs	2.4	0.02483165
	AB002340 at	M33493	tryptase, beta (tryptase II)16p13.3	2.4	0.02689938
	RC_AA446661 at	AB002340	KIAA0342 gene product	2.3	0.000748796
	RC_AA446661_at RC_AA084138_at	AA446661	ESTs	2.3	0.011980248
	NO_AA004138_at	AA084138	ESTs	2.3	1.16025E-05
	DC N50966 -1	NEOGGO	ESTs, Weakly similar to putative		0.000
	RC_N59866_at	N59866	p150 [H.sapiens]	2.3	0.002042263
	RC_R42424_at	R42424	ESTs	2.3	0.003173074
	RC_N39415_at	N39415	osteoglycin (osteoinductive factor)	2.3	0.001310764

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	2 vs BPH-With Sympt		TABLE 1		
1654533.1	A.EE. I	Genbank	Genbank	Fold-change	p-value
	Affy element	ID IO0404	Name	N1-N2 vs With	
	J03464_s_at	J03464	collagen, type I, alpha 27q22.1	2.3	0.006791534
	RC_AA205376_at	AA205376	KIAA0471 gene product1q24-q25	2.3	0.023123837
	RC H95960 at	H95960	secreted protein, acidic, cysteine-rich	2.3	0.008509182
	NG_H95900_at	парабо	(osteonectin)5q31.3-q32 bone marrow stromal cell antigen	2.3	0.000509102
	D28137_at	D28137	219p13.2	2.3	0.031127266
	D20107_at	D20101	extracellular matrix protein 2, female	2.0	0.001127200
	RC_N79778 at	N79778	organ and adipocyte specific9q22.3	2.3	0.045073744
	RC_N98485_s_at	N98485	forkhead (Drosophila)-like 66p25.3	2.3	0.033372862
			prostaglandin D2 synthase (21kD,	2.0	0,0000.2002
	M98539_at	M98539	brain)9q34.2-q34.3	2.2	0.005442674
	RC_AA205724_at	AA205724	ESTs	2.2	0.006183612
			Homo sapiens ribonuclease 6		
	U85625_at	U85625	precursor, mRNA, complete cds.	2.2	0.001245066
			RAB2, member RAS oncogene		
	RC_R37588_s_at	R37588	family-like6p21.3	2.2	0.00219386
	RC_AA046426_at	AA046426	Cdc42 effector protein 3	2.2	0.005788723
	RC_AA256294_at	AA256294	ESTs	2.2	0.002425605
			SWI/SNF related, matrix associated,		
	DO 11700100 1	4.4.500.4.50	actin dependent regulator of		0.0100=0011
	RC_AA599120_at	AA599120	chromatin, subfamily e, member 1	2.2	0.042979241
	RC_W60186_at	W60186	ESTs	2.2	0.028494835
	RC AA599216 at	AA599216	collapsin response mediator protein 14p16.1-p15	2.2	0.040523744
	RC AA450324 at	AA450324	ESTs	2.2	0.040523744
	110_701+3002+_at	AA400024	Homo sapiens aldehyde	2.1	0.009094307
	M31994 at	M31994	dehydrogenase (ALDH1) gene	2.1	0.001561218
	RC_AA402930 at	AA402930	ESTs	2.1	0.000114627
		,	Human AMP deaminase isoform L		0.000111021
			(AMPD2) mRNA, exons 6-18, partial		
	M91029_cds2_at	M91029_cds2	cds	2.1	0.02494373
		_	ESTs, Weakly similar to 17beta-		
			hydroxysteroid dehydrogenase		
	RC_AA450114_at	AA450114	[H.sapiens]	2.1	4.87556E-06
	D62584_at	D62584	osteoglycin (osteoinductive factor)	2.1	0.000157116
	RC_AA621634_at	AA621634	ESTs	2.1	0.02297009
	RC_AA312946_s_at	AA312946	ESTs	2.1	3.51075E-05
	V07429 a at	V07420	Human DNA for cellular retinol	0.4	0.000045047
	X07438_s_at	X07438	binding protein (CRBP)	2.1	0.039015947
	RC_N53447_at	N53447	integral membrane protein 2CXq21.1 21.2	2.1	0.009032297
	110_1100++1_at	14004-17	Homo sapiens mRNA; cDNA	2.1	0.009032291
			DKFZp586B211 (from clone		
	RC_AA281591 at	AA281591	DKFZp586B211)	2.0	0.016660714
			ESTs, Moderately similar to	2.0	0.010000111
			alternatively spliced product using		
	RC_R71395_at	R71395	exon 13A [H.sapiens]	2.0	0.046231847
			cytochrome P450, subfamily XIA		
			(cholesterol side chain		
	RC_T53590_s_at	T53590	cleavage)15q23-q24	2.0	0.00282074
	RC_AA293489_at	AA293489	KIAA0638 protein	2.0	0.006966532
	RC_AA447707_s_at	AA447707	KIAA1055 protein	2.0	0.001248537
	RC_AA235618_f_at	AA235618	ESTs	2.0	0.012481746
	RC_N68350_at	N68350	ESTs	2.0	0.035156598
	RC_H81379_s_at	LI04270	ESTs, Moderately similar to	2.0	0.04440400
	110_11013/9_5_at	H81379	KIAA0438 [H.sapiens]	2.0	0.01148429
	RC_D51060_s_at	D51060	Jun activation domain binding	2.0	0.046668054
	D01000_s_at	201000	protein1p32-p31 B-cell translocation gene 2	2.0	0.016668951
	U72649_at	U72649	(pheochromacytoma cell-3)1q32	2.0	0.020660388
	RC_AA287389_at	AA287389	ESTs	2.0	0.002741873
	RC_AA621367_at	AA621367	ESTs	2.0	0.004871903
			secreted protein, acidic, cysteine-rich		
	J03040_at	J03040	(osteonectin)5q31.3-q32	2.0	0.006303994
			• • •		

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	al2 vs BPH-With Symp		TABLE 1	Cald shames	
1654533.1	Affy element	Genbank ID	Genbank Name	Fold-change	p-value N1-N2 vs With
	Any element	ID	non-metastatic cells 5, protein	NI-NZ VS VVIUI	IN I-INZ VS VVIUI
			expressed in (nucleoside-		
	RC_AA291676_s_at	AA291676	diphosphate kinase)5q23-q31	2.0	0.027480479
	RC_N63536_at	N63536	ESTs	2.0	0.000634305
			UDP-Gal:betaGlcNAc beta 1,3-		
			galactosyltransferase, polypeptide		
	RC_AA411952_at	AA411952	33q25	2.0	
	RC_AA252802_s_at RC_AA382275_at	AA252802 AA382275	Human mRNA for TI-227H ESTs	2.0 2.0	
	NC_AA302213_at	AA302213	tissue inhibitor of metalloproteinase	2.0	0.00087437
	AA093923 at	AA093923	217g25	2.0	0.046200886
	M11313 s at	M11313	alpha-2-macroglobulin12p13.3-p12.3	2.0	
	RC_AA398280_at	AA398280	ESTs	2.0	
	RC_N51529_at	N51529	ESTs	2.0	0.006276979
			nudix (nucleoside diphosphate linked		
	H49440_at	H49440	moiety X)-type motif 36p21.2	2.0	
	RC_T33263_s_at	T33263	KIAA0320 protein	2.0	
	RC_T89160_r_at	T89160	ESTs Weekly similar to	2.0	0.005289266
			ESTs, Weakly similar to serine/threonine protein kinase TAO1		
	RC W56792 at	W56792	[R.norvegicus]	2.0	0.026130523
			ESTs, Moderately similar to	2.0	0.020100020
			alternatively spliced product using		
	RC_R60056_at	R60056	exon 13A [H.sapiens]	2.0	0.001585076
			Human Chromosome 16 BAC clone		
Down-regulated		AA398908	CIT987SK-A-61E3	-21.7	
	RC_AA460914_at RC_T40895 at	AA460914	ESTs	-15.8	
	NO_140095_at	T40895	ESTs ESTs, Moderately similar to FAT-	-12.6	0.002430219
			SPECIFIC PROTEIN FSP27		
	RC_R71792_s_at	R71792	[M.musculus]	-9.8	0.01438632
	RC_N80129_i_at	N80129	metallothionein 1L16q13	-8.7	
			myosin, light polypeptide 2,		
	X66141_at	X66141	regulatory, cardiac, slow12q23-q24.3	-8.0	0.03928942
	A A O D 4 C O 4 . F k	4.400.400.4	CCAAT/enhancer binding protein		
	AA234634_f_at	AA234634	(C/EBP), delta8p11.2-p11.1	-7.4	0.000589696
	U78294 at	U78294	arachidonate 15-lipoxygenase, second type	-6.8	0.017271608
	RC_AA457566_at	AA457566	ESTs	-6.6	
		, , , , , , , , , , , , , , , , , , , ,	phospholemman-like, expressed in	-0.0	0.023044022
	X93036_at	X93036	breast tumors, 8kD	-6.2	0.011323909
	X57129_at	X57129	H1 histone family, member 26p21.3	-6.1	0.004161922
			Human intestinal mucin mRNA,		
	HG1067-HT1067_r_at		partial cds, clone SMUC 42	-5.8	0.007202185
	X65614_at	X65614	S100 calcium-binding protein P4p16	-5.8	0.006892572
	RC_AA609006_at	AA609006	ESTs	-5.7	0.015701354
	J03910_rna1_at RC H94471 at	J03910_rna1 H94471	metallothionein 1G16q13 occludin5q13.1	-5.7	0.003506953
	AB000584 at	AB000584	prostate differentiation factor	-5.6 -5.4	0.025014274 0.003235425
	RC_W88568 at	W88568	glycogenin 2Xp22.3	-5.4 -5.1	0.048573115
	V00594_at	V00594	metallothionein 2A16q13	-5.0	0.000721258
	RC_T73433_s_at	T73433	angiotensinogen1q41-qter	-4.9	0.012700144
	RC_N94303_at	N94303	ESTs	-4.5	4.88059E-05
			Homo sapiens mRNA; cDNA		
	DO 4440044 /		DKFZp586D0823 (from clone		
	RC_AA419011_at	AA419011	DKFZp586D0823)	-4.1	0.013801595
	RC_N32748_at	N32748	ESTs Weakly similar to much Muc?	-4.1	0.018749207
	RC_AA053424 at	AA053424	ESTs, Weakly similar to mucin Muc3 [R.norvegicus]	-4.0	0.001235197
	RC_AA599331_at	AA599331	ESTs	-4.0 -4.0	0.001233197
			folate hydrolase (prostate-specific		5.555.5550
	M99487_at	M99487	membrane antigen) 111p11.2	-3.9	0.013268152
	RC_F02245_at	F02245	monoamine oxidase AXp11.4-p11.3	-3.8	0.002950391
		X76717	metallothionein 1L16q13	-3.7	0.000868707
	X64177_f_at	X64177	metallothionein 1H16q13	-3.7	0.002089771

Man 14 M			33		
	al2 vs BPH-With Symp	toms rabie Genbank	TABLE 1	Cold observe	
1654533.1	ı Affy element	ID	Genbank	Fold-change N1-N2 vs With	p-value
	Any element	טו	Name squamous cell carcinoma antigen	14 1-142 VS VVIIII	IN 1-INZ VS VVILII
	RC_AA599522_r_at	AA599522	recognised by T cells	-3.6	0.012643918
			human homolog of yeast		
	L77701 at	L77701	mitochondrial copper recruitment	-3.6	0.002244007
	LITTUI_at	LITTOI	gene	-3.0	0.003341007
			ESTs, Moderately similar to weak		
	RC D11824 at	D11824	similarity to Arabidopsis thaliana ubiquitin-like protein 8 [C.elegans]	-3.6	0.000803294
	RC_AA410311 at	AA410311	ESTs	-3.5	0.001234064
	RC AA457235 at	AA457235	ESTs	-3.5	0.012177965
			protein tyrosine phosphatase type	5.0	0.0.2.7.000
	RC_N93798_at	N93798	IVA, member 3	-3.5	0.007340453
			nuclear receptor subfamily 1, group		
	RC_AA416762_s_at	AA416762	H, member 219q13.3-19q13.3	-3.5	0.010404304
			ESTs, Weakly similar to tumorous		
			imaginal discs protein Tid56 homolog		
	RC_F03969_at	F03969	[H.sapiens]	-3.5	0.011826812
	RC_AA045487_at	AA045487	ESTs	-3.4	0.025187615
	RC_Z38744_at	Z38744	putative gene product13	-3.4	2.30674E-05
			ESTs, Moderately similar to HERV-E		
	RC_N92502_s_at	N92502	integrase [H.sapiens]	-3.4	0.02301359
	RC_R91484_at	R91484	ESTs	-3.4	8.2306E-05
	RC_AA165313_at	AA165313	ESTs	-3.3	0.028364404
	RC_AA182030_at	AA182030	ESTs	-3.3	0.019770486
			ESTs, Moderately similar to (defline		
	RC_T94447_s_at	T94447	not available 4335935) [M.musculus]	-3.3	0.001427294
	RC_W20486_f_at	W20486	ESTs	-3.3	0.002892697
	RC_R16983_at	R16983	ESTs	-3.2	0.000912559
			interferon stimulated gene		
	RC_AA504805_s_at	AA504805	(20kD)15q26	-3.2	0.003905701
	RC_T90190_s_at	T90190	H1 histone family, member 26p21.3	-3.2	0.020618793
	RC_AA135870_at	AA135870	ESTs	-3.1	0.04609197
	RC_H99035_at	H99035	ESTs	-3.1	0.000191451
	RC_R28370_at	R28370	ESTs	-3.1	0.024606319
			alcohol dehydrogenase 3 (class I),		
	RC_T40995_f_at	T40995	gamma polypeptide4q21-q23	-3.1	0.024064044
	MIP1-B_at	MIP1-B	karyopherin (importin) beta 2	-3.1	0.005882353
			ESTs, Highly similar to differentially		
	20 1111		expressed in Fanconi anemia		
	RC_AA447522_at	AA447522	[H.sapiens]	-3.1	0.003518059
	DO 444644		ESTs, Moderately similar to Cab45a		
	RC_AA461453_at	AA461453	[M.musculus]	-3.0	0.021949087
	AA429539_f_at	AA429539	ESTs	-3.0	0.017623102
	RC_AA476944_at	AA476944	ESTs	-3.0	0.019974254
	RC_N80129_f_at	N80129	metallothionein 1L16q13	-3.0	0.000219038
			ESTs, Weakly similar to		
	DC NOCOOA	NOCOOA	FK506/rapamycin-binding protein		
	RC_N26904_at	N26904	FKBP13 precursor [H.sapiens]	-2.9	0.006305062
	RC_AA505136_at	AA505136	ESTs	-2.9	0.005400284
	AA455001_s_at	AA455001	ESTs	-2.9	2.1534E-05
	RC_W70131_at	W70131	ESTs	-2.9	0.005764635
	RC_AA043349_at	AA043349	ESTs	-2.9	0.016983419
	U02020_at	U02020	pre-B-cell colony-enhancing factor	-2.9	0.003324497
	U52969_at	U52969	Purkinje cell protein 421q22.2-q22.3	-2.8	0.00078638
	RC_H22453_at	H22453	ESTs	-2.8	0.000410695
	RC_N22620_at RC N64683 at	N22620	ESTs	-2.8	0.005507089
		N64683	ESTs	-2.8	0.00378977
	RC_N24761_at RC_AA464728_s_at	N24761	ESTs	-2.8	0.004837185
	RC_AA464728_s_at RC_H83380_at	AA464728 H83380	ESTs ESTs	-2.8 -2.7	0.004669897
	1.0_1100000_at	1100000		-2.1	0.016543793
	M30894_at	M30894	T-cell receptor, gamma cluster7p15- p14	-2.7	0.004450467
	RC_H81070 f at	H81070	P14 Human metallothionein (MT)I-F gene	-2.7 -2.7	0.034153167
	a.	1.01010	actin, alpha, cardiac muscle15q11-	-2.1	0.022654931
	J00073_at	J00073	gter	-2.7	0.029724167
			401	-2.1	0.028/2410/

Normal1-Normal2 vs BPH-V	Vith Symptoms Tal	34 Die TABLE 1		
1654533.1	Genban	k Genbank	Fold-change	p-value
Affy elemen	nt ID	Name	N1-N2 vs With	N1-N2 vs With
		ESTs, Weakly similar to ORF		
RC_H05084	4_at H05084		-2.7	0.016965435
		Homo sapiens mRNA; cDNA		
		DKFZp564A072 (from clone		
AA045870_	at AA0458		-2.7	0.005480167
RC_T68873	3_f_at T68873	metallothionein 1L16q13	-2.7	0.001140431
RC_N72253	3_at N72253		-2.7	0.001832591
		Homo sapiens mRNA; cDNA		
		DKFZp564A072 (from clone		
RC_AA4479			-2.7	0.001255304
RC_H18947		ESTs	-2.7	0.00193501
RC_H77597		metallothionein 1H16q13	-2.7	0.001560766
RC_H94475			2 -2.6	0.01435663
RC_AA0253	370_at AA0253	70 KIAA0872 protein	-2.6	0.013924142
		ESTs, Moderately similar to PIM-1		
		PROTO-ONCOGENE		
		SERINE/THREONINE-PROTEIN		
RC_AA4431	_	14 KINASE [M.musculus]	-2.6	0.000703574
RC_F09684	_	ESTs	-2.6	0.000107291
RC_AA0313	360_s_at AA0313	60 ESTs	-2.6	0.047293081
RC_AA4166	685_at AA4166	(-: -:-gane) intecp :   p : 22	-2.6	0.023296279
		UDP-Gal:betaGlcNAc beta 1,4-		
		galactosyltransferase, polypeptide		
D29805_at	D29805	19p13	-2.6	2.3562E-05
		solute carrier family 2 (facilitated		
		glucose transporter), member 11p	35-	
RC_H58873	3_s_at H58873	p31.3	-2.5	0.000710917
M10942_at	M10942	metallothionein 1E (functional)16q	13 -2.5	0.017370635
RC_T03593	3_at T03593	ESTs	-2.5	0.006239127
		small inducible cytokine A5		
RC_N95495	5_at N95495	(RANTES)17q11.2-q12	-2.5	0.002392984
		ESTs, Highly similar to Miz-1 prote	eln	
RC_AA0170		63 [H.sapiens]	-2.5	0.048093776
RC_R00144	1_at R00144	ESTs	-2.5	0.018222161
		squamous cell carcinoma antigen		
RC_AA5995			-2.5	
RC_AA2195	552_s_at AA2195		-2.5	0.043156485
		ESTs, Moderately similar to (deflin		
RC_AA4475	_	in a constant of the constant		
RC_AA0707	752_s_at AA0707		-2.5	0.002895462
DQ D00000		ESTs, Weakly similar to cappuccin		_
RC_R02003		[D.melanogaster]	-2.4	
L13698_at	L13698	growth arrest-specific 19q21.3-q22	.1 -2.4	0.013393145
DC 444000	000 -4 0000	ESTs, Moderately similar to B cell		
RC_AA4322	292_at AA4322	State of the state	-2.4	0.000956642
•		DNA segment, single copy probe		
RC_H99648	) o of 100649	LNS-CAI/LNS-CAII (deleted in	0.4	0.0000000
RC AA1319		polyposis5q22-q23	-2.4	
RC_AA6216	_	_ ,	-2.4	
110	695_at AA62169		-2.4	0.008761556
		ESTs, Weakly similar to !!!! ALU		
RC AA5986	695 at AA5986	SUBFAMILY SX WARNING ENTR		0.000540077
NC_AA3900	195_at AA5900	[r i.oapicrio]	-2.4	0.000549977
		ESTs, Moderately similar to !!!! AL		
RC_AA4303	888 at AA4303	SUBFAMILY SQ WARNING ENTR		0.000405470
M24069_at	M24069	[eaptorie]	-2.4 -2.4 -2.4	
W2+009_at	1012-4003	cold shock domain protein A12p13	.1 -2.4	0.015890231
RC_AA4341	08 at AA43410	Homo sapiens heat shock protein	-2.4	0.040400000
RC_AA4054		map is a military complete and	-2.4 -2.3	
RC_AA4195	_	· - · -	-2.3 -2.3	
RC_W38197		EST	-2.3 -2.3	
1.0_1.00197	7700107	superoxide dismutase 2,	-2.3	0.013006462
RC_R38709	_s_at R38709	mitochondrial6q25.3	-2.3	0.02567404
1.0_1.00100		ESTs, Moderately similar to coppe		0.03567491
RC_AA1211	42 at AA12114	transport protein HAH1 [H.sapiens]		0.043639016
1.0_1.1.211		adioport protein HART [In.saplens]	2.0	U.U43U38U10

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Normal1-Normal2 vs BPH-With Symp		TABLE 1		
1654533.1	Genbank	Genbank		value
Affy element	ID	Name	N1-N2 vs With N	1-N2 vs With
RC_N26801_at	N26801	ESTs	-2.3	0.000580867
RC_N75960_at	N75960	ESTs	-2.3	0.01244791
RC_R36969_at	R36969	ESTs	-2.3	0.019129486
	1100000		-2.0	0.013123400
AA046840 at	AA046840	CCAAT/enhancer binding protein (C/EBP), delta8p11.2-p11.1	-2.3	0.002504544
7.1.0.100.10_0.1	7 2 10 100 10	transforming, acidic coiled-coil	2.0	0.002001011
RC R46074 at	R46074	containing protein 210q26	-2.3	0.003462273
X06956 at	X06956	tubulin, alpha 1 (testis specific)2q	-2.3	0.015437809
RC_H84761_s_at	H84761	glutathione peroxidase 13p21.3	-2.2	0.000365528
RC_W52065_f_at	W52065		-2.2	
110_1102000_1_at	VV32003	KIAA0539 gene product	-2.2	0.016497348
RC_AA279757_at	AA279757	ESTs, Weakly similar to (defline not available 4481810) [D.melanogaster]	-2.2	0.003272622
110_7 11270701_at	701270707	ESTs, Weakly similar to (defline not	-2.2	0.003272022
RC_H16676_s_at	H16676	available 5107634) [R.norvegicus]	-2.2	8.86866E-05
RC AA255480 at	AA255480	ESTs	-2.2	
				0.009359024
RC_R96924_s_at	R96924	ESTs	-2.2	0.000201685
		ESTs, Moderately similar to !!!! ALU		
•		SUBFAMILY SQ WARNING ENTRY		
RC_AA342337_at	AA342337	!!!! [H.sapiens]	-2.2	0.024999347
RC_AA004699_at	AA004699	putative translation initiation factor	-2.2	0.022298405
		tumor suppressor deleted in oral		
RC_AA401965_at	AA401965	cancer-related 111g13	-2.2	0.006294885
		Homo sapiens clone 24796 mRNA		
RC_F02470_at	F02470	sequence	-2.2	0.022313149
	. 02110	sodium channel, nonvoltage-gated 1	-2.2	0.022010149
X76180 at	X76180	, ,	0.0	0.000070004
7/0100_at	X10100	alpha12p13	-2.2	0.023078001
DC D40400	D40400	coatomer protein complex, subunit		
RC_R49138_s_at	R49138	epsilon	-2.2	0.020401578
		actin related protein 2/3 complex,		
RC_D80237_s_at	D80237	subunit 4 (20 kD)	-2.2	0.022022634
		growth arrest and DNA-damage-		
RC_AA402224_at	AA402224	inducible, gamma9q22.1-q22.2	-2.2	0.014983528
		Homo sapiens mRNA for for histone		
RC_AA281599_at	AA281599	H2B, clone piG4-5-14	-2.2	0.029567009
RC_N78630_at	N78630	KIAA0870 protein	-2.2	0.006668895
X85785_rna1_at	X85785 rna1	•		
	_	Duffy blood group1q21-q22	-2.2	0.018706507
RC_AA412063_at	AA412063	ESTs	-2.2	0.000686563
		ESTs, Weakly similar to		
		phosphatidylinositol transfer protein		
RC_AA022886_at	AA022886	[H.sapiens]	-2.2	0.000777067
RC_N24899_at	N24899	ESTs	-2.2	0.030610964
RC_AA101767_at	AA101767	ESTs	-2.2	0.009040467
		ESTs, Weakly similar to Homo		
RC_AA045503_at	AA045503	sapiens p20 protein [H.sapiens]	-2.2	0.021950966
RC F10078 at	F10078	ESTs	-2.1	0.040699115
RC_H02308_at	H02308			
		ESTs	-2.1	0.036730715
RC_AA284153_at	AA284153	ESTs	-2.1	0.021270233
RC_AA453433_at	AA453433	HLA-B associated transcript-16p21.3	-2.1	0.013366375
		Homo sapiens Ste-20 related kinase		
RC_AA403159_at	AA403159	SPAK mRNA, complete cds	-2.1	0.025212073
		Homo sapiens clone 23836 mRNA		
RC_T17428_s_at	T17428	sequence	-2.1	0.044754602
		ESTs, Highly similar to (defline not		
RC_W92449 at	W92449	available 4587714) [H.sapiens]	-2.1	0.019386585
RC_AA609312_at	AA609312	ESTs	-2.1	0.003204911
. 10_7 0 10000 12_41	000012		-2.1	J.UUJZU4811
D28589 at	D28589	Human mRNA (KIAA00167), partial	0.4	0.000400470
D20003_at	220008	sequence	-2.1	0.000408478
DC AAGGEOG at	V V 333500	ESTs, Highly similar to (defline not		0.0010000
RC_AA232508_at	AA232508	available 4929647) [H.sapiens]	-2.1	0.004626663
RC_AA280929_s_at	AA280929	ESTs	-2.1	0.028189798
	11/00====	S-adenosylmethionine decarboxylase		
W63793_at	W63793	16q21-q22	-2.1	0.032076011
		Homo sapiens DNA from		
		chromosome 19-cosmid R30879		
RC_R36881_s_at	R36881	containing USF2, genomic sequence	-2.1	0.007343473
RC_AA278767_s_at	AA278767	ESTs	-2.1	0.001983494
		· <del>-</del>		2.00 1000 10-F

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	al2 vs BPH-With Symp		TABLE 1		
1654533.1		Genbank	Genbank	Fold-change	p-value
	Affy element	ID	Name		N1-N2 vs With
	RC_R98442_at	R98442	ESTs	-2.1	
	X99728_at	X99728	H.sapiens NDUFV3 gene, exon 3.	-2.1	0.001404191
			solute carrier family 11 (proton-		
	DO DOCCES /		coupled divalent metal ion		
	RC_R09379_at	R09379	transporters), member 212q13	-2.1	0.006004344
	RC R99092 at	R99092	EST, Moderately similar to (defline not available 5052951) [H.sapiens]	-2.1	0.016256526
	X95325 s at	X95325	cold shock domain protein A12p13.1	-2.1	0.025953179
	RC_T56281 f_at	T56281	Human metallothionein (MT)I-F gene	-2.1	0.032089569
	RC R44397 at	R44397	ESTs	-2.1	
	RC_H27180_f_at	H27180	ESTs	-2.1	0.004317675
	AA165312 at	AA165312	ESTs	-2.1	0.025559572
	RC_AA279313_s_at	AA279313	methyl CpG binding protein 2Xq28	-2.1	0.030594523
			Homo sapiens beta-tubulin mRNA,		0.00000.000
	HG4322-HT4592_at	AF141349	complete cds.	-2.1	0.017120749
	-		high-mobility group (nonhistone		
			chromosomal) protein isoforms I and		
	RC H81413 f at	H81413	Y6p21	-2.1	0.009976588
			ESTs, Highly similar to (defline not		0,0000,0000
	RC_W94333 at	W94333	available 5107163) [H.sapiens]	-2.1	0.000435688
			eukaryotic translation initiation factor		0.000.0000
	RC AA455070 at	AA455070	3, subunit 1 (alpha, 35kD)	-2.1	0.025226928
	RC_R11526_f_at	R11526	parathymosin17q12-q22	-2.1	0.027182202
	RC T15409 f at	T15409	EST EST	-2.1	0.001478856
	RC_H05625_f_at	H05625	ESTs	-2.1	0.024564209
	RC_AA620461_at	AA620461	ESTs	-2.0	
	RC_AA449791_f at	AA449791	EST	-2.0 -2.0	
	RC AA435769 s at	AA435769	ESTs	-2.0	
	RC N55502 at	N55502	ESTs ·	-2.0	
	110_1100002_01	1100002	tumor suppressing subtransferable	-2.0	0.021034433
	AF001294_at	AF001294	candidate 311p15.5	-2.0	0.03566128
			ESTs, Highly similar to (defline not	2.0	0.00000120
	RC Z40898 at	Z40898	available 4929639) [H.sapiens]	-2.0	0.002289892
	RC_AA436861_at	AA436861	ESTs	-2.0	0.00187676
	7147 7110000 7		peptidylprolyl isomerase B	2.0	0.00101010
	M63573 at	M63573	(cyclophilin B)15	-2.0	0.044239663
	RC_T25732 f_at	T25732	KIAA0252 protein	-2.0	
			ESTs, Weakly similar to (defline not		313 1 1 2 1 3 3 3
	RC R01257 at	R01257	available 4456991) [H.sapiens]	-2.0	0.005735841
	RC_H91703_i_at	H91703	cell division cycle 2717q12-17q23.2	-2.0	
	RC N34817 at	N34817	ESTs	-2.0	0.040996591
			ESTs, Weakly similar to KIAA0374		
	RC_R60777_at	R60777	[H.sapiens]	-2.0	0.000245565
			ESTs, Weakly similar to		
			MICROTUBULE-ASSOCIATED		
	RC_AA386264_at	AA386264	PROTEIN 1B [M.musculus]	-2.0	0.000541139
			ESTs, Weakly similar to Containing		
			ATP/GTP-binding site motif A(P-		
			loop): Similar to C.elegans		
			protein(P1:CEC47E128);Similar to		
			Mouse alpha-		
			mannosidase(P1:B54407)		
	RC_AA251769_at	AA251769	[H.sapiens]	-2.0	0.008985897
	RC_R56602_at	R56602	lg superfamily proteinXq12-q13.3	-2.0	0.024051216
	RC_AA397919_at	AA397919	ESTs	-2.0	
			ESTs, Weakly similar to envelope		
	RC_W37778_f_at	W37778	protein [H.sapiens]	-2.0	0.043013942
	AA248555_at	AA248555	ESTs	-2.0	0.000824698
	,		ESTs, Weakly similar to		
			SERINE/THREONINE-PROTEIN		
	RC_AA463693_at	AA463693	KINASE NEK3 [H.sapiens]	-2.0	0.002809026
	M70404 1	W70404	NADH dehydrogenase (ubiquinone) 1	_	0.4
	W76181_at	W76181	alpha subcomplex, 2 (8kD, B8)5q31	-2.0	0.008370263
	RC_AA171939_at	AA171939	ESTs	-2.0	0.015796116

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Normal1-Norn	nal2 vs BPH-With Symp	toms Table	TABLE 1		
1654533		Genbank	Genbank	Fold-change	p-value
	Affy element	ID	Name	N1-N2 vs With	N1-N2 vs With
			U30999 Homo sapiens MV3		
			melanoma Homo sapiens cDNA		
	U30999_at	U30999	clone memd	-2.0	0.007070546
			synuclein, alpha (non A4 component		
	RC_F03254_f_at	F03254	of amyloid precursor)4q21	-2.0	0.011479379
			ESTs, Weakly similar to !!!! ALU		
			SUBFAMILY SC WARNING ENTRY		
	RC_H26288_at	H26288	!!!! [H.sapiens]	-2.0	
	RC_AA007158_f_at	AA007158	ESTs	-2.0	0.001870921
		•	Homo sapiens clone 23940 mRNA		
	RC_Z38785_at	Z38785	sequence	-2.0	
	RC_AA282247_at	AA282247	ESTs	-2.0	0.000515617
			ESTs, Weakly similar to protein-		
	RC_T23935_s_at	T23935	tyrosine phosphatase [H.sapiens]	-2.0	
·	RC_R59593_at	R59593	ESTs	-2.0	
	RC_AA446241_at	AA446241	tropomyosin 2 (beta)9p13.2-p13.1	-2.0	0.040680667
			DJ222E13.1a.1 (C-terminal part of		
			novel protein dJ222E13.1) (partial		
	RC_Z40556_at	Z40556	isoform 1)	-2.0	0.019444878
			ESTs, Highly similar to (defline not		
	RC_AA159025_at	AA159025	available 4680655) [H.sapiens]	-2.0	0.01375696
			estrogen-responsive B box		
	RC_H03387_s_at	H03387	protein17p11.2	-2.0	
	RC_H17333_at	H17333	EST	-2.0	0.018111182
			putative cyclin G1 interacting		
	RC_AA412722_s_at	AA412722	protein7	-2.0	0.006838915
			NADH dehydrogenase (ubiquinone)		
			Fe-S protein 8 (23kD) (NADH-		
	U65579_at	U65579	coenzyme Q reductase)11q13	-2.0	
	RC_R88209_at	R88209	ESTs	-2.0	0.040272012
	D.O. #200000	=	Homo sapiens PAC clone		0.000444000
	RC_Z38266_at	Z38266	DJ0777O23 from 7p14-p15	-2.0	0.009414008

		30				
Normal1-Normal2 vs BPH-Cancer Table	Up- regulated	Table 2				
1654552.1	Genbank	Genbank			Fold-Change	p-value
Affy element	ID	Name			N1-N2 vs	N1-N2 vs
					Cancer	Cancer
L49169_at	L49169	FBJ murine osteosarcoma vira	al oncogene hom	olog B19q13.3	18.8	0.03580379
RC_N23730_s_at	N23730	v-fos FBJ murine oste homolog14q24.3	eosarcoma vira	al oncogene	16.5	8.98673E-05
V01512_rna1_at	V01512_rn a1		eosarcoma vir	al oncogene	16.0	0.001216643
RC_T90619_f_at	T90619	actin, gamma 117q25			15.7	0.044124187
U20734_s_at	U20734	jun B proto-oncogene19p13.2			14.3	0.004404553
U62015_at	U62015	insulin-like growth factor bindir	ng protein 101p2:	2-p <b>31</b>	13.8	0.000487216
AA374109_at	AA374109	ESTs, Moderately similar to ([R.norvegicus]	defline not avail	able 5031506)	13.0	0.025911461
RC_T79768_at	T79768	ESTs			12.2	0.018940142
RC_AA410383_at	AA410383	B-cell-homing chemokine (li receptor-1)4a21	gand for Burki	tt's lymphoma	11.1	0.046025784
X52541_at	X52541	early growth response 15q31.	I		9.7	0.003167537
RC_N66802_at	N66802	early growth response 38p23-	021		9.7	0.026764792
RC_AA463726_s_at	AA463726	JM27 proteinXp11.23			9.4	0.003409168
N40141_at	N40141	JM27 proteinXp11.23			8.4	0.021768214
M34996_s_at	M34996	major histocompatibility compl	ex, class II, DQ a	alpha 16p21.3	7.7	0.015886207
RC_T67053_f_at	T67053	immumoglobulin lambda gene	cluster22q11.1-0	11.2	7.4	0.000196865
RC_AA404957_at	AA404957	ESTs, Highly similar to PRECURSOR [H.sapiens]	MATRIX (	GLA-PROTEIN	6.6	0.011451385
RC_H64493_f_at	H64493	immunoglobulin gamma 3 (Gr	n marker)14q32.3	33	6.5	0.002716347
RC_N47686_s_at	N47686	solute carrier family 14 (urea blood group)18q11-q12	transporter), m	ember 1 (Kidd	6.3	0.015568892
RC_W44760_s_at	W44760	frizzled-related protein2qter			6.3	0.016891036
L19871_at	L19871	activating transcription factor 3	3		6.2	0.007603286
M92934_at	M92934	connective tissue growth factor	r6q23.1		6.1	0.001046931
M62831_at	M62831	immediate early protein19			5.8	0.00753286
L22524_s_at	L22524	matrix metalloproteinase 7 (ma	atrilysin, uterine)	11q21-q22	5.8	0.048289798
J03507_at	J03507	complement component 75p13	3		5.6	0.00240657
RC_AA236455_r_at	AA236455	ESTs			5.5	0.022653542
RC_AA450127_at	AA450127	growth arrest and DNA-damag	e-inducible, beta	19p13.3	5.5	0.023227588
RC_AA281345_f_at	AA281345	immediate early protein19			5.4	0.003661068
RC_N30198_at	N30198	ESTs			5.3	0.005657756
AFFX-	X00351	Human mRNA for beta-actin			5.3	0.01547291
HSAC07/X00351_5_a t	l .					
D83018_at	D83018	nel (chicken)-like 212q13.11-q	13.12		5.1	0.003774757
J04111_at	J04111	Jun activation domain binding	protein1p32-p31		5.0	0.000243067
X51345_at	X51345	jun B proto-oncogene19p13.2			5.0	0.017173421
RC_AA398903_at	AA398903	ESTs, Weakly similar to !!!! A ENTRY !!!! [H.sapiens]	ALU SUBFAMILY	/ J WARNING	4.9	0.014577818
RC_H17550_at	H17550	ESTs			4.7	0.012079391
S81914_at	S81914	immediate early response 36p	21.3		4.5	0.006218653
RC_AA250958_f_at	AA250958	EST			4.4	1.88343E-05
RC_AA446651_at	AA446651	ESTs			4.4	0.026022802
HG1872-HT1907_at	M28590	Human (clone pcDG-79) MH partial cds.	C HLA-DG prote	ein 41 mRNA,	4.3	0.008830524
RC_AA490667_at	AA490667	ESTs			4.3	0.048863016
RC_N67041_at	N67041	ESTs			4.1	0.009333688
V00563_at	V00563	immunoglobulin mu14q32.33			4.1	0.004301939
X57809_s_at	X57809	immumoglobulin lambda gene	cluster22q11.1-0	11.2	4.1	0.025371658

Normal1-Normal2 vs BPH-Cancer Table	Up- regulated	Table 2		
1654552,1	Genbank	Genbank	Fold-Change	p-value
Affy element	ID	Name	N1-N2 vs Cancer	N1-N2 vs Cancer
R69417_at	R69417	ESTs	4.1	0.046373179
J00231_f_at	J00231	immunoglobulin gamma 3 (Gm marker)14q32.33	4.0	0.004766015
RC_AA402903_f_at	AA402903	immunoglobulin gamma 3 (Gm marker)14q32.33		
U21128_at	U21128	lumican12q21.3-q22	3.9	0.000172905
M12529_at	M12529		3.9	0.000708917
RC_AA436616_at	AA436616	apolipoprotein E19q13.2 ESTs	3.7	0.026856247
U72649_at	U72649		3.7	0.020860083
X03689_s_at		B-cell translocation gene 2 (pheochromacytoma cell-3)1q32	3.7	0.002487396
	X03689	Human mRNA fragment for elongation factor TU (N-terminus)	3.7	0.04821902
AFFX- HSAC07/X00351_5_a	X00351	Human mRNA for beta-actin	3.6	0.029717275
t RC_T62857_at	T62857	ESTs	3.6	0.000046530
Z74616_s_at	Z74616			0.002846539
X06700_s_at		collagen, type II, alpha 1 (Ehlara Daylos gyndrama type IV	3.6	0.004328291
	X06700	collagen, type III, alpha 1 (Ehlers-Danlos syndrome type IV, autosomal dominant)2q31	3.6	0.010596098
RC_H86112_f_at	H86112	KIAA0471 gene product1q24-q25	3.6	0.017013968
M57466_s_at	M57466	major histocompatibility complex, class II, DP beta 16p21.3	3.5	0.005924671
RC_F09281_at	F09281	ESTs	3.5	0.006841731
RC_R51831_at	R51831	ESTs	3.4	0.000941423
RC_H21814_f_at	H21814	immumoglobulin lambda gene cluster22q11.1-q11.2	3.4	0.009767098
RC_W86513_at	W86513	ESTs	3.4	0.003776481
RC_H40424_s_at	H40424	EST	3.4	0.016283906
X57025_at	X57025	insulin-like growth factor 1 (somatomedin C)12q22-q23	3.3	0.040489253
RC_AA044219_at	AA044219	BK984G1.1 (PUTATIVE C-terminal end of a novel protein with Collagen triple helix repeats)	3.3	0.001761114
RC_AA028092_s_at	AA028092	transcription factor 216pter-qter	3.3	0.003405482
RC_AA446661_at	AA446661	ESTs	3.3	0.041188995
RC_D80063_f_at	D80063	ESTs	3.3	0.049585142
M92843_s_at	M92843	zinc finger protein homologous to Zfp-36 in mouse19q13.1	3.3	0.006174082
M34516_r_at	M34516	immunoglobulin lambda-like polypeptide 322q11.2	3.2	0.02344053
M87789_s_at	M87789	immunoglobulin gamma 3 (Gm marker)14q32.33	3.2	0.004534646
N75870_s_at	N75870	dual specificity phosphatase 15q34	3.2	0.000157434
RC_AA609309_at	AA609309	ESTs, Moderately similar to !!!! ALU SUBFAMILY SB2 WARNING ENTRY !!!! [H.sapiens]	3.1	0.03780658
S59049_at	S59049	regulator of G-protein signalling 11q31	3.0	0.002419303
AFFX-	M33197	Human GAPDH	3.0	0.034538288
HUMGAPDH/M33197 5 at				0.00
RC_D51060_s_at	D51060	Jun activation domain binding protein1p32-p31	3.0	0.022390037
RC_T23468_at	T23468	ESTs	2.9	0.001634616
U30521_at	U30521	P311 protein	2.9	0.009484198
Z48501_s_at	Z48501	poly(A)-binding protein-like 13q22-q25	2.9	0.026396977
W73859_at	W73859	transcription factor 216pter-qter	2.9	0.037326183
AA093923_at	AA093923	tissue inhibitor of metalloproteinase 217q25	2.8	0.041564022
RC_AA236476_at	AA236476	ESTs, Weakly similar to (defline not available 4507549)		0.038305276
U10550_at	U10550	[H.sapiens] GTP-binding protein overexpressed in skeletal muscle8q13-	2.7	0.040657885
RC_N24902_at	N24902	q21 E1B-55kDa-associated protein 5	2.7	0.03810507
RC_AA056121_at	AA056121	ESTs	2.7	0.03810307
				0.027200100

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Normal1-Normal2 vs BPH-Cancer Table	Up- regulated	Table 2		
1654552.1	Genbank	Genbank	Fold-Change	p-value
Affy element	ID	Name	N1-N2 vs Cancer	N1-N2 vs Cancer
RC_H98835_at	H98835	ESTs	2.7	0.019901442
K02405_f_at	K02405	Human MHC class II HLA-DQ-beta mRNA (DR7 DQw2), complete cds	2.7	0.00138806
U90552_s_at	U90552	butyrophilin, subfamily 3, member A16p23	2.7	3.91186E-05
RC_N59831_at	N59831	ESTs	2.7	0.04543669
L33799_at	L33799	procollagen C-endopeptidase enhancer7q22	2.7	0.010879277
RC_N59532_s_at	N59532	aminomethyltransferase (glycine cleavage system protein T)3p21.2-p21.1	2.6	0.025712285
D13628_at	D13628	anglopoletin 18q22.3-q23	2.6	0.027204836
AA156897_s_at	AA156897	Homo sapiens mRNA; cDNA DKFZp564I1922 (from clone DKFZp564I1922)	2.6	0.001580022
RC_N67876_s_at	N67876	insulin-like growth factor 1 (somatomedin C)12q22-q23	2.6	0.03992641
M73720_at	M73720	carboxypeptidase A3 (mast cell)3q21-q25	2.6	0.023298997
H49440_at	H49440	nudix (nucleoside diphosphate linked moiety X)-type motif 36p21.2	2.6	0.002498701
RC_AA250850_at	AA250850	adrenergic, beta, receptor kinase 222q11	2.5	0.041156086
RC_T49061_at	T49061	ESTs	2.5	0.00934004
W28214_at	W28214	ESTs	2.5	0.037677921
RC_H44631_s_at	H44631	immediate early protein19	2.5	0.0423037
D28137_at	D28137	bone marrow stromal cell antigen 219p13.2	2.5	0.026212334
RC_AA609027_at	AA609027	ESTs	2.5	0.038550623
RC_AA257093_r_at	AA257093	T-cell receptor, beta cluster7q35	2.4	0.002653232
RC_F13763_at	F13763	ESTs	2.4	0.016949277
RC_H08548_s_at	H08548	ATP citrate lyase17q12-q21	2.4	0.036998522
RC_AA436618_at	AA436618	ESTs	2.4	0.001789907
RC_W45664_s_at	W45664	5' nucleotidase (CD73)6q14-q21	2.4	0.001762727
AA082546_at	AA082546	ESTs	2.4	0.021791878
D10522_at	D10522	myristoylated alanine-rich protein kinase C substrate (MARCKS, 80K-L)6q22.2	2.4	0.017333686
RC_AA411860_at	AA411860	ESTs, Highly similar to (defline not available 4929723) [H.saplens]	2.4	0.02766922
AB002340_at	AB002340	KIAA0342 gene product	2.3	0.003238699
U53445_at	U53445	downregulated in ovarian cancer 13	2.3	0.009361652
AA091278_at	AA091278	ESTs	2.3	0.046253689
RC_AA486072_i_at	AA486072	small inducible cytokine A5 (RANTES)17q11.2-q12	2.3	0.012816473
RC_T53590_s_at	T53590	cytochrome P450, subfamily XIA (cholesterol side chain cleavage)15q23-q24	2.3	4.29636E-05
RC_N91971_f_at	N91971	retinol-binding protein 1, cellular3q23	2.3	0.025171598
RC_AA043777_at	AA043777	ESTs	2.3	0.004490188
RC_H54764_at	H54764	EST, Weakly similar to X-linked retinopathy protein {C-terminal, clone XEH.8c} [H.sapiens]	2.3	0.036980431
RC_AA443923_at	AA443923	ESTs	2.3	0.025833241
U60975_at	U60975	Homo sapiens gp250 precursor, mRNA, complete cds.	2.3	0.041238204
M34516_at	M34516	immunoglobulin lambda-like polypeptide 322q11.2	2.3	0.041388637
RC_N36001_at	N36001	ESTs, Weakly similar to !!!! ALU CLASS C WARNING ENTRY !!!! [H.sapiens]	2.2	0.000449076
AF010193_at	AF010193	MAD (mothers against decapentaplegic, Drosophila) homolog 718	2.2	0.005397771
AFFX- HSAC07/X00351_5_a t	X00351	Human mRNA for beta-actin	2.2	0.037852217
RC_AA158262_s_at	AA158262	calpastatin5q14-q22	2.2	0.006648962
RC_AA156565_at	AA156565	4-nitrophenylphosphatase domain and non-neuronal SNAP25 like 122q12	2.2	0.020901922

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Normal1-Normal2 vs BPH-Cancer Table	Up- regulated	Table 2		
1654552.1	Genbank	Genbank	Fold-Change	p-value
Affy element	ID	Name .	N1-N2 vs	N1-N2 vs
			Cancer	Cancer
Z11793_at	Z11793	selenoprotein P, plasma, 15q31	2.2	0.00118281
RC_D80059_s_at	D80059	ESTs	2.2	0.033534432
RC_AA450324_at	AA450324	ESTs	2.2	0.024832006
RC_N39415_at	N39415	osteoglycin (osteoinductive factor)	2.2	0.032001116
RC_T23622_at	T23622	ESTs	2.2	0.040417825
RC_AA599365_at	AA599365	decorin12q23	2.2	0.011325181
X62320_at	X62320	granulin17	2.2	0.043043858
RC_R85291_at	R85291	ESTs	2.2	0.004987693
M11313_s_at	M11313	alpha-2-macroglobulin12p13.3-p12.3	2.2	0.011545737
AA047151_at	AA047151	ESTs	2.2	0.033987576
RC_AA205724_at	AA205724	ESTs	2.2	0.004569368
RC_AA086264_i_at	AA086264	ESTs, Highly similar to (defline not available 4191348)	2.2	0.020637423
RC_R42424_at	R42424	[H.sapiens] ESTs	2.2	0.033603417
RC_AA347359_s_at	AA347359	lysozyme (renal amyloidosis)12	2.1	0.028764499
AA092716_at	AA092716	HLA-B associated transcript-36p21.3	2.1	0.031717351
RC_R42241_at	R42241	ESTs	2.1	0.008013968
RC_N57577_at	N57577	KIAA0663 gene product	2.1	0.032028875
RC_W67577_s_at	W67577	CD74 antigen (invariant polypeptide of major histocompatibility complex, class II antigen-associated)5q32	2.1	0.002072118
C02016_at	C02016	KIAA0447 gene product	2.1	0.002399894
RC_AA256268_at	AA256268	ESTs	2.1	0.0269568
RC_T96171_at	T96171	EST	2.1	0.012219229
X72841_at	X72841	retinoblastoma-binding protein 7	2.1	0.033774692
RC_R45698_at	R45698	ESTs	2.1	0.049975895
RC_N22006_s_at	N22006	EST	2.1	0.011131338
RC_N69222_at	N69222	ESTs	2.1	0.022256915
RC_H97538_at	H97538	ESTs	2.0	0.03795259
RC_AA039935_at	AA039935	dynein light chain, outer arm 422q12.3-q13.2	2.0	0.011488766
RC_AA084138_at	AA084138	ESTs	2.0	0.011124432
AB002379_at	AB002379	KIAA0381 protein	2.0	0.000530413
RC_AA460651_at	AA460651	heterogeneous nuclear protein similar to rat helix destabilizing protein10	2.0	0.027697892
RC_W02204_at	W02204	solute carrier family 24 (sodium/potassium/calcium exchanger), member 115q22	2.0	0.00115779
Y08614_at	Y08614	exportin 1 (CRM1, yeast, homolog)2p16	2.0	0.035368368
D31134_at	D31134	KIAA1075 protein	2.0	0.021196526
M94880_f_at	M94880	major histocompatibility complex, class I, A6p21.3	2.0	0.025382167
J03040_at	J03040	secreted protein, acidic, cysteine-rich (osteonectin)5q31.3-q32	2.0	0.035472553
RC_N68350_at	N68350	ESTs	2.0	0.042917893
RC_H48793_at	H48793	EST	2.0	0.00296551
HG3543-HT3739_at	M29645	insulin-like growth factor 2 (somatomedin A)11p15.5	2.0	0.019712374
RC_W33172_at	W33172	ESTs, Weakly similar to ORF2 [M.musculus]	2.0	0.006454106
RC_R08850_at	R08850	ESTs	2.0	0.011364766
W52638_at	W52638	ESTs	2.0	0.010612401
M19045_f_at	M19045	lysozyme (renal amyloidosis)12	2.0	0.004561974
RC_AA312946_s_at	AA312946	ESTs	2.0	0.020272205
RC_AA235310_at	AA235310	ESTS	2.0	0.011954937
X03100_cds2_at	X03100_cd s2	Human mRNA for SB classII histocompatibility antigen alphachain	2.0	0.002404541

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Normal1-Normal2 vs BPH-Cancer Table	Up- regulated	Table 2		
1654552.1	Genbank	Genbank	Fold-Change	p-value
Affy element	ID	Name	N1-N2 vs	N1-N2 vs Cancer
RC_T16282_f_at	T16282	wood+ (\$ namba) hamalagddad5 2 nd5 d	Cancer	
RC_H66642_f_at	H66642	wee1+ (S. pombe) homolog11p15.3-p15.1 ESTs, Moderately similar to !!!! ALU SUBFAMILY SQ	2.0	0.031472155
NC_1100042_1_at	F100042	WARNING ENTRY !!!! [H.sapiens]	2.0	0.02460529
RC_AA342337_at	AA342337	ESTs, Moderately similar to !!!! ALU SUBFAMILY SQ WARNING ENTRY !!!! [H.sapiens]	-23.7	3.26344E-05
RC_AA398908_at	AA398908	Human Chromosome 16 BAC clone CIT987SK-A-61E3	-21.7	0.040053626
RC_H15143_s_at	H15143	Human clone 23575 mRNA, partial cds	-13.8	0.028261625
RC_N80129_i_at	N80129	metallothionein 1L16q13	-12.6	0.002146038
RC_AA465394_at	AA465394	ESTs	-12.6	0.004961162
RC_AA236545_at	AA236545	ESTs	-12.5	0.034938167
RC_W42778_at	W42778	Homo sapiens clone 24636 mRNA sequence	-12.3	0.010449419
RC_T40895_at	T40895	ESTs	-12.0	0.01968535
RC_H94475_s_at	H94475	alpha-2-plasmin inhibitor17pter-p12	-11.7	0.012919819
RC_R71792_s_at	R71792	ESTs, Moderately similar to FAT-SPECIFIC PROTEIN FSP27 [M.musculus]	-10.4	0.002540356
RC_AA609006_at	AA609006	ESTs	-7.5	0.013902978
RC_AA026641_s_at	AA026641	secretory leukocyte protease inhibitor (antileukoproteinase)	-7.0	0.01850877
X65614_at	X65614	S100 calcium-binding protein P4p16	-6.7	0.005634308
X93036_at	X93036	phospholemman-like, expressed in breast tumors, 8kD	-6.6	0.005278275
RC_T94447_s_at	T94447	ESTs, Moderately similar to (defline not available 4335935) [M.musculus]	-5.7	0.006891909
RC_AA405488_at	AA405488	ESTs	-5.5	0.00023986
RC_T73433_s_at	T73433	angiotensinogen1q41-qter	-5.5	0.009418205
M99487_at	M99487	folate hydrolase (prostate-specific membrane antigen) 111p11.2	-5.3	0.008067789
RC_W88568_at	W88568	glycogenin 2Xp22.3	-5.1	0.024739084
RC_AA460914_at	AA460914	ESTs	-5.0	0.024385552
X57129_at	X57129	H1 histone family, member 26p21.3	-4.8	0.006322499
RC_Z41642_at	Z41642	ESTs	-4.7	0.009525521
RC_R46074_at	R46074	transforming, acidic coiled-coil containing protein 210q26	-4.7	0.001327844
J03910_rna1_at	J03910_rna 1	metallothionein 1G16q13	-4.6	0.004574277
RC_AA350265_at	AA350265	histone deacetylase A	-4.5	0.002897414
AA165312_at	AA165312	ESTs	-4.2	0.005487803
RC_AA419011_at	AA419011	Homo sapiens mRNA; cDNA DKFZp586D0823 (from clone DKFZp586D0823)	-4.0	0.019079557
RC_N92502_s_at	N92502	ESTs, Moderately similar to HERV-E integrase [H.sapiens]	-4.0	0.030144039
RC_F03969_at	F03969	ESTs, Weakly similar to tumorous imaginal discs protein Tid56 homolog [H.sapiens]	-4.0	0.017024613
X76717_at	X76717	metallothionein 1L16q13	-3.9	0.001145402
RC_AA416762_s_at	AA416762	nuclear receptor subfamily 1, group H, member 219q13.3-19q13.3	-3.8	0.011735303
RC_AA053424_at	AA053424	ESTs, Weakly similar to mucin Muc3 [R.norvegicus]	-3.8	0.009737433
X64177_f_at	X64177	metallothionein 1H16q13	-3.7	0.003297195
RC_N32748_at	N32748	ESTS	-3.6	0.021454174
RC_AA416685_at	AA416685	UNC13 (C. elegans)-like9p11-p12	-3.6	0.016338392
RC_AA505136_at	AA505136	ESTS	-3.5	0.007200396
RC_AA165313_at	AA165313	ESTS	-3.5	0.037649191
RC_F02245_at	F02245	monoamine oxidase AXp11.4-p11.3	-3.4	0.005486135
RC_AA004699_at	AA004699	putative translation initiation factor	-3.4	0.00057505
RC_AA599331_at	AA599331	ESTs	-3.4	0.01136457

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Normal1-Normal2 vs BPH-Cancer Table	Up- regulated	Table 2	E-14 Ch	
1654552.1	Genbank	Genbank	Fold-Change	p-value
Affy element	ID	Name	N1-N2 vs Cancer	N1-N2 vs Cancer
RC_N26904_at	N26904	ESTs, Weakly similar to FK506/rapamycin-binding protein FKBP13 precursor [H.sapiens]	-3.3	0.045410608
RC_AA070752_s_at	AA070752	insulin receptor substrate 12q36	-3.3	0.028433761
RC_AA599522_f_at	AA599522	squamous cell carcinoma antigen recognised by T cells	-3.2	0.005311305
RC_N94303_at	N94303	ESTs	-3.1	0.000160723
RC_F10078_at	F10078	ESTs	-3.1	0.022464594
RC_AA447537_at	AA447537	ESTs, Moderately similar to (defline not available 5360237) [M.musculus]	-3.1	0.007323728
L77701_at	L77701	human homolog of yeast mitochondrial copper recruitment	-3.0	0.001489928
RC_H27675_at	H27675	gene ESTs	-3.0	0.016160504
V00594_at	V00594	metallothionein 2A16q13	-2.9	0.001495259
U52969_at	U52969	Purkinje cell protein 421q22.2-q22.3	-2.9	6.3447E-05
RC_R42607_at	R42607	ESTs	-2.8	0.008960052
RC_AA451836_at	AA451836	ESTs	-2.7	0.008401586
RC_F04492_at	F04492	ESTs, Weakly similar to !!!! ALU SUBFAMILY J WARNING ENTRY !!!! [H.sapiens]	-2.7	0.001443051
RC_H77597_f_at	H77597	metallothionein 1H16q13	-2.7	0.00332868
RC_AA430388_at	AA430388	ESTs, Moderately similar to !!!! ALU SUBFAMILY SQ	-2.7	0.000114004
RC_T90190_s_at	T90190	WARNING ENTRY !!!! [H.sapiens] H1 histone family, member 26p21.3	-2.7	0.030242714
RC_H16171_f_at	H16171	cleft lip and palate associated transmembrane protein	-2.7	0.023414443
RC_AA022886_at	AA022886	119q13.2-q13.3 ESTs, Weakly similar to phosphatidylinositol transfer protein [H.sapiens]	-2.7	0.00489294
RC_R28370_at	R28370	ESTs	-2.7	0.003724547
RC_AA261907_at	AA261907	ESTs, Weakly similar to (defline not available 3874144) [C.elegans]	-2.6	0.043689441
RC_W37778_f_at	W37778	ESTs, Weakly similar to envelope protein [H.sapiens]	-2.6	0.030756837
RC_T98019_at	T98019	EST, Highly similar to PEREGRIN [H.sapiens]	-2.5	0.035566681
RC_N33927_s_at	N33927	H2B histone family, member B6p21.3	-2.5	0.013093926
RC_R40431_at	R40431	Homo sapiens mRNA; cDNA DKFZp564D016 (from clone DKFZp564D016)	-2.5	0.004235538
RC_AA133756_at	AA133756	Rho-associated, coiled-coil containing protein kinase 22p24	-2.5	0.012389163
RC_AA152200_s_at	AA152200	ESTs	-2.5	0.004366137
W63793_at	W63793	S-adenosylmethionine decarboxylase 16q21-q22	-2.5	0.005714247
RC_AA410298_at	AA410298	ESTs	-2.5	0.018744617
X99728_at	X99728	H.sapiens NDUFV3 gene, exon 3	-2.5	0.004580383
RC_W78127_at	W78127	ESTs, Weakly similar to KIAA0425 [H.sapiens]	-2.5	0.001240164
RC_R96924_s_at	R96924	ESTs	-2.5	0.006515911
RC_H16768_at	H16768	ESTs	-2.5	0.005669237
X76180_at	X76180	sodium channel, nonvoltage-gated 1 alpha12p13	-2.5	0.007625025
RC_AA432162_at	AA432162	Homo sapiens mRNA; cDNA DKFZp586B2022 (from clone DKFZp586B2022)	-2.4	0.010199113
RC_H88798_at	H88798	ESTs	-2.4	0.000783143
RC_AA609312_at	AA609312	ESTs	-2.4	0.016243321
RC_AA131919_at	AA131919	putative type II membrane protein	-2.4	0.000264791
RC_N80129_f_at	N80129	metallothionein 1L16q13	-2.4	0.002297016
RC_AA182030_at	AA182030	ESTs	-2.4	0.041632378
W70167_at	W70167	ESTs	-2.4	0.00395969
RC_AA599522_r_at	AA599522	squamous cell carcinoma antigen recognised by T cells	-2.4	0.004347078
RC_N52254_s_at	N52254	SH3-binding domain glutamic acid-rich protein21q22.3	-2.4	0.011171389
RC_N95495_at	N95495	small inducible cytokine A5 (RANTES)17q11.2-q12	-2.4	0.002430242

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Normal1-Normal2 vs BPH-Cancer Table	Up- regulated	Table 2		
1654552.1	Genbank	Genbank	Fold-Change	p-value
Affy element	ID	Name	N1-N2 vs	N1-N2 vs
			Cancer	Cancer
RC_T68873_f_at	T68873	metallothionein 1L16q13	-2.4	0.00320019
AA429539_f_at	AA429539	ESTs	-2.4	0.020751882
RC_AA435769_s_at	AA435769	ESTs	-2.4	0.009832353
RC_AA029356_at	AA029356	ESTs	-2.3	0.007208722
AA316686_s_at RC_H02308_at	AA316686 H02308	ESTs, Highly similar to huntingtin interacting protein HYPK [H.sapiens] ESTs		0.000225753
RC_AA258476_at	AA258476	Homo sapiens mRNA; cDNA DKFZp564J0323 (from clone	-2.3 -2.3	0.041776289
X06956 at	X06956	DKFZp564J0323) tubulin, alpha 1 (testis specific)2q	-2.3 -2.3	0.02070961 0.003656874
RC H99694 at	H99694	ESTs	-2.3 -2.3	0.003636874
RC_AA479044_s_at	AA479044	ESTs, Weakly similar to PROGASTRICSIN PRECURSOR	-2.3 -2.3	0.047032301
1.0_111110011_0_at	, , , , , , , , , , , , , , , , , , , ,	[H.sapiens]	-2.0	0.047 03230 1
RC_AA436861_at	AA436861	ESTS	-2.3	0.001794201
M24069_at	M24069	cold shock domain protein A12p13.1	-2.3	0.014123514
RC_AA410311_at	AA410311	ESTs	-2.3	0.045227011
W52858_at	W52858	Homo sapiens mRNA; cDNA DKFZp564F0522 (from clone DKFZp564F0522)	-2.3	0.002276405
RC_W38197_at	W38197	EST	-2.3	1.96016E-05
J00073_at	J00073	actin, alpha, cardiac muscle15q11-qter	-2.3	0.018476889
RC_D51069_f_at	D51069	melanoma adhesion molecule	-2.3	0.042693395
RC_AA504805_s_at	AA504805	interferon stimulated gene (20kD)15q26	-2.3	0.008805886
RC_F03254_f_at	F03254	synuclein, alpha (non A4 component of amyloid precursor)4q21	-2.3	0.003668915
M35252_at	M35252	transmembrane 4 superfamily member 3	-2.3	0.028083185
RC_AA040731_at	AA040731	ESTs	-2.2	0.028924808
RC_AA496247_at	AA496247	ESTs	-2.2	0.013336314
X59766_at	X59766	alpha-2-glycoprotein 1, zinc7	-2.2	0.002003511
RC_R84421_at	R84421	eukaryotic translation elongation factor 1 alpha 16q14	-2.2	0.016333706
AA328993_s_at	AA328993	ESTs	-2.2	0.004438605
RC_R44535_f_at	R44535	endonuclease G9q34.1	-2.2	0.014319616
U41518_at	U41518	aquaporin 1 (channel-forming integral protein, 28kD)7p14	-2.2	0.009447457
RC_W33179_at	W33179	testis-specific kinase 21p32	-2.2	0.001104272
RC_H58873_s_at	H58873	solute carrier family 2 (facilitated glucose transporter), member 11p35-p31.3	-2.2	0.000238641
RC_R31679_s_at	R31679	ESTS	-2.2	0.01000414
RC_AA189083_at	AA189083	ESTs, Highly similar to (defline not available 4589468) [M.musculus]	-2.2	0.002468046
RC_AA251769_at	AA251769	ESTs, Weakly similar to Containing ATP/GTP-binding site motif A(P-loop): Similar to C.elegans protein(P1:CEC47E128);Similar to Mouse alphamannosidase(P1:B54407) [H.sapiens]	-2.2	0.010819016
RC_W70131_at	W70131	ESTs	-2.2	0.02955725
RC_R09379_at	R09379	solute carrier family 11 (proton-coupled divalent metal ion	-2.2	0.009730513
RC AA621695 at	AA621695	transporters), member 212q13	0.4	0.004004004
RC_H18947_at	H18947	ESTs · · · · · · · · · · · · · · · · · · ·	-2.1	0.001994051
			-2.1	0.027246274
RC_AA219552_s_at	AA219552	ESTS	-2.1	0.046510941
RC_N22620_at	N22620	ESTs Weakly similar to conquesing ID malay a sector	-2.1	0.013527392
RC_R02003_r_at	R02003	ESTs, Weakly similar to cappuccino [D.melanogaster]	-2.1 ·	0.010597095
RC_AA405559_at	AA405559	ESTS	-2.1	0.009305601
RC_AA463693_at	AA463693	ESTs, Weakly similar to SERINE/THREONINE-PROTEIN KINASE NEK3 [H.sapiens]	-2.1	0.004156996

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Normal1-Normal2 vs BPH-Cancer Table	Up- regulated	Table 2		
1654552.1	Genbank	Genbank	Fold-Change	p-value
Affy element	ID	Name	N1-N2 vs	N1-N2 vs
			Cancer	Cancer
RC_AA481407_at	AA481407	ESTs	-2.1	0.002741696
M11119_at	M11119	Human endogenous retrovirus envelope region mRNA (PL1)	-2.1	0.003718876
RC_AA159025_at	AA159025	ESTs, Highly similar to (defline not available 4680655) [H.sapiens]	-2.1	0.011127532
RC_AA411981_at	AA411981	ESTs, Weakly similar to putative seven pass transmembrane protein [H.sapiens]	-2.1	0.044294612
RC_W57931_at	W57931	ESTs, Moderately similar to CATHEPSIN D PRECURSOR [H.sapiens]	-2.1	0.000755739
X66899_at	X66899	Ewing sarcoma breakpoint region 122q12	-2.1	0.002068901
RC_R49327_at	R49327	solute carrier family 11 (proton-coupled divalent metal ion transporters), member 212q13	-2.1	0.030928835
RC_AA609645_at	AA609645	eukaryotic translation initiation factor 4 gamma, 13q27-qter	-2.1	0.04955957
RC_AA434108_at	AA434108	Homo sapiens heat shock protein hsp40-3 mRNA, complete cds	-2.1	0.034468752
X17567_s_at	X17567	small nuclear ribonucleoprotein polypeptides B and B120	-2.1	0.014475221
J04164_at	J04164	interferon-induced protein 17	-2.1	0.023410352
RC_AA135929_s_at	AA135929	ESTs, Highly similar to (defline not available 4103057) [M.musculus]	-2.1	0.003009065
L04270_at	L04270	lymphotoxin beta receptor (TNFR superfamily, member 312p13	-2.1	0.006776988
RC_H99035_at	H99035	ESTs	-2.1	0.001053884
M64673_at	M64673	heat shock transcription factor 1	-2.1	0.004283001
X85785_rna1_at	X85785_rn a1	Duffy blood group1q21-q22	-2.1	0.00657464
M68864_at	M68864	Human ORF mRNA, complete cds	-2.1	0.010185833
D50928_at	D50928	KIAA0138 gene product	-2.1	0.002283064
RC_AA282247_at	AA282247	ESTs	-2.0	0.007970044
RC_R00144_at	R00144	ESTs	-2.0	0.006939854
RC_AA485965_at	AA485965	ESTs, Highly similar to (defline not available 4336766) [H.sapiens]	-2.0	0.000405037
S45630_at	S45630	crystallin, alpha B11q22.3-q23.1	-2.0	0.006157273
RC_T89703_at	T89703	ESTs, Highly similar to (defline not available 4455129) [H.sapiens]	-2.0	0.000286616
RC_Z38785_at	Z38785	Homo sapiens clone 23940 mRNA sequence	-2.0	0.00706437
X85373_at	X85373	small nuclear ribonucleoprotein polypeptide G	-2.0	6.93881E-05
RC_F04816_at	F04816	ESTs	-2.0	0.005353184
RC_AA043349_at	AA043349	ESTs	-2.0	0.01749596
RC_H84761_s_at	H84761	glutathione peroxidase 13p21.3	-2.0	0.000116621
M34338_s_at	M34338	spermidine synthase1p36-p22	-2.0	0.008566137
L13698_at	L13698	growth arrest-specific 19q21.3-q22.1	-2.0	0.016504513
RC_N75960_at	N75960	ESTs	-2.0	0.024082428
D45370_at	D45370	adipose specific 210	-2.0	0.034362163
RC_AA401965_at	AA401965	tumor suppressor deleted in oral cancer-related 111q13	-2.0	0.011190087
RC_F09315_at	F09315	discs, large (Drosophila) homolog 510q23	-2.0	0.020753036
RC_AA025370_at	AA025370	KIAA0872 protein	-2.0	0.026565555
RC_H52835_at	H52835	phytanoyl-CoA hydroxylase (Refsum disease)10pter-p11.2	-2.0	0.015021251
RC_H99648_s_at	H99648	DNA segment, single copy probe LNS-CAI/LNS-CAII (deleted in polyposis5q22-q23	-2.0	0.012115852
RC_AA430074_at	AA430074	ESTs		0.002355049
RC_AA598939_at	AA598939	ESTs		0.011383872
AA455001_s_at	AA455001	ESTs		0.000176199
RC_F09684_at	F09684	ESTs		0.002741682
D42073_at	D42073	reticulocalbin 1, EF-hand calcium binding domain11p13	-2.0	0.012881688

Normal1-Normal2 vs BPH-Cancer Table	Up- regulated	Table 2		
1654552.1	Genbank	Genbank	Fold-Change	p-value
Affy element	ID	Name	N1-N2 vs	N1-N2 vs
			Cancer	Cancer
RC_AA598695_at	AA598695	ESTs, Weakly similar to !!!! ALU SUBFAMILY SX WARNING ENTRY !!!! [H.sapiens]	-2.0	4.77268E-06
D23662_at	D23662	neural precursor cell expressed, developmentally down- regulated 8	-2.0	0.003156141
RC_AA431470_at	AA431470	protein kinase (cAMP-dependent, catalytic) inhibitor gamma20g	-2.0	0.038692982
RC_AA399273_at	AA399273	ESTs	-2.0	0.029403118
RC_AA142858_at	AA142858	ESTs	-2.0	0.00197166
RC_Z40715_at	Z40715	Homo sapiens mRNA; cDNA DKFZp586C201 (from clone DKFZp586C201)	-2.0	0.017206338
RC_AA490341_s_at	AA490341	ESTs	-2.0	0.004570941
RC_N67815_f_at	N67815	ESTs, Weakly similar to (defline not available 4680655) [H.sapiens]	-2.0	0.002996692
RC_N53359_at	N53359	ESTs	-2.0	0.034916164

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	W/Symptoms Table		GenBank Name	Fold-change	t
Up-regulated	-		JM27 protein	Ü	
1	_	N40141	v-fos FBJ murine osteosarcoma viral oncogene	17.4	-7.64
2 3		N23730	5M29 Stein	10.8 10.0	-7.54
3 4	rc_AA463726_s_at rc_N23352 s at	N23352	proenkephalin	10.0	-6.56
5		H64493	immunoglobulin heavy constant gamma 3 (G3m	9.1	-4.53 -4.36
6		V01512	marker) FBJ murine osteosarcoma viral oncogene	9.1	-7.40
7		H05704	here (29helix coiled-coil rod homologue)	8.1	-2.79
8		L49169	FBJ murine osteosarcoma viral oncogene homolog B	8.0	-2.79 -5.81
	_		B-cell-homing chemokine (ligand for Burkitt's lymphoma		
9		AA410383	receptor-1)	7.5	-3.95
10	rc_AA131322_s_at	AA131322	tryptase, alpha,tryptase, beta (tryptase II) eukaryotic translation initiation factor 3, subunit 6 (48kD)	7.2	-2.81
11	R56183_s_at	R56183		6.9	-2.77
12	rc_AA461300_at	AA461300	ESTs	6.9	-7.08
13	J00231_f_at	J00231	immunoglobulin heavy constant gamma 3 (G3m	6.7	-4.62
14	rc_AA427622_s_at	AA427622	ଓଖିଣ୍ଡିନେ, type XIII, alpha 1	6.6	-8.25
15		T90889	ESTs	5.6	-3.72
16	rc_AA402903_f_at		immunoglobulin heavy constant gamma 3 (G3m	5.6	-3.61
17	_ <u>-</u>	T23622	matter)	5.5	-5.24
18		T62857	ESTs	5.4	-7.85
19		AA256268	ESTs	5.3	-6.86
20		R44714	ESTs	5.3	-4.83
21		AA236476	transmembrane protein TENB2,	5.1	-3.13
22	rc_AA028092_s_at		transcription factor 21	5.1	-5.24
23		T90619	actin, gamma 1	5.0	-2.19
24	_	J00123	proenkephalin	5.0	-3.96
25	_	X52541	early growth response 1	4.9	-5.78
26		AA620825	CGI-43 protein ESTs	4.9	-4.59
27	rc_AA424530_s_at	AA424530	procollagen-proline, 2-oxoglutarate 4-dioxygenase	4.9	-5.42
28	rc_AA386386_s_at	AA386386	(proline 4-hydroxylase), beta polypeptide (protein disulfide isomerase; thyroid hormone binding protein p55)	4.9	-2.64
29	U62015_at	U62015	cysteine-rich, angiogenic inducer, 61	4.9	-6.24
30	rc_AA188981_at	AA188981	highly expressed in cancer, rich in leucine heptad repeats	4.9	-6.67
31	rc_H21814_f_at	H21814	immunoglobulin lambda locus	4.9	-2.67
32	M60314_at	M60314	bone morphogenetic protein 5		-10.82
33	rc_T67053_f_at	T67053	immunoglobulin lambda locus	4.7	-2.84
34	rc_N47686_s_at	N47686	solute carrier family 14 (urea transporter), member 1 (Kidd blood group)	4.7	-3.27
35	rc_AA436616_at	AA436616	ESTs	4.7	-6.34
36	rc_H60595_s_at	H60595	progesterone binding protein	4.7	-2.66
37	rc_H88338_at	H88338	ESTs	4.7	-7.93
38	M33653_at		collagen, type XIII, alpha 1	4.6	-8.95
39			ESTs	4.5	-5.87
40			nel (chicken)-like 2	4.5	-9.79
41			ESTs ( Table 1)	4.5	-6.27
42		H61295	CD4 antigen (p55)	4.4	-4.49
43	rc_AA281345_f_at .		immediate early protein	4.3	-6.62
			hypothetical protein FLJ20185	4.2	-5.25
45	rc_AA279760_at	AA279760	DKFZP564M182 protein	4.2	-3.73

Normal vs. BPH W/Symptoms Table **TABLE 3** 1654540.1

Normal vs. BPH	W/Symptoms Table	TABLE 3	1654540.1		
Up-regulated	Affy element	GenBank ID	GenBank Name	Fold-change	t
46	rc_R25410_at	R25410	ESTs	4.2	-4.69
47	rc_T03229_f_at	T03229	ESTs	4.2	-3.37
48	rc_R93908_at	R93908	ESTs	4.2	-3.39
49	AA374109_at	AA374109	spondin 2, extracellular matrix protein	4.2	-1.97
50	rc_R45654_at	R45654	collagen, type XIII, alpha 1	4.2	-5.69
51	rc_H86112_f_at	H86112	KIAA0471 gene product	4.1	-4.00
52	rc_AA257093_r_at	AA257093	T cell receptor beta locus	4.1	-7.77
53	rc_AA456147_at	AA456147	general transcription factor IIIA	4.1	-6.23
54	U21128_at	U21128	lumican	4.1	-6.15
55	rc_AA057195_at	AA057195	TNF? elastin microfibril interface located protein	4.1	-2.22
56	M63438_s_at	M63438	immunoglobulin kappa variable 1D-8	4.0	-2.53
57	M57466_s_at	M57466	major histocompatibility complex, class II, DP beta 1	4.0	-3.91
58	rc_AA443923_at	AA443923	cat eye syndrome critical region gene 1	4.0	-3.01
59	rc_N39415_at	N39415	DKFZP586P2421 protein	4.0	-5.70
60	rc_W67225_at	W67225	KIAA0592 protein	4.0	-3.35
61	M62831_at	M62831	immediate early protein	4.0	-6.39
62	rc_AA404957_at	AA404957	matrix Gla protein	4.0	-3.84
63	rc_F02992_at	F02992	ESTs	4.0	-3.65
64	U69263_at	U69263	matrilin 2	3.9	-4.84
65	rc_AA448625_at	AA448625	slit (Drosophila) homolog 3	3.9	-4.13
66	X57025_at	X57025	insulin-like growth factor 1 (somatomedin C)	3.9	-3.93
67	AA151544_at	AA151544	matrix metalloproteinase 23B	3.8	-5.54
68	rc_F13763_at	F13763	ESTs	3.8	-6.39
69	rc_AA436655_at	AA436655	hypothetical protein FLJ10781	3.8	-5.13
70	M87789_s_at	M87789	immunoglobulin heavy constant gamma 3 (G3m		-3.93
71	L44416_at	L44416	DEAGNH (Asp-Glu-Ala-Asp/His) box polypeptide 17 (72kD)		-1.75
72	U20350_at	U20350	chemokine (C-X3-C) receptor 1	3.8	-6.50
73	rc_AA449749_at	AA449749	ESTs	3.8	-4.52
74	rc_W73790_f_at	W73790	immunoglobulin lambda-like polypeptide 1	3.7	-2.95
75	rc_AA281145_at	AA281145	ESTs	3.7	-1.77
76	rc_f09748_s_at	f09748	ESTs	3.7	-4.12
77	rc_T64211_at	T64211	HNOEL-iso protein	3.7	-5.35
78	rc_N80152_at	N80152	RNA binding motif protein 6	3.7	-2.40
79	rc_AA436618_at	AA436618	microtubule-associated protein 2	3.7	-4.67
80	T85532_f_at	T85532	ESTs	3.7	-1.90
81	rc_AA398280_at	AA398280	ESTs ·	3.6	-3.11
82	rc_T23468_at	T23468	CGI-119 protein	3.6	-4.67
83	AA195678_at	AA195678	actin binding protein; macrophin (microfilament and actin filament cross-linker protein)	3.6	-3.48
84	AB002335_at	AB002335	KIAA0337 gene product	3.6	-4.21
85	rc_AA598982_s_at	AA598982	KIAA1114 protein,trophinin	3.6	-4.58
86	J03507_at	J03507	complement component 7	3.6	-6.21
87	J04130_s_at	J04130	small inducible cytokine A4 (homologous to mouse Mip- 1b)	3.5	-4.76
88	AA495865_at	AA495865	ESTs	3.5	-3.65
89	HG3543-HT3739_		insulin-like growth factor 2 (somatomedin A)	3.5	-4.69
90	rc_AA599662_s_at		KIAA0534 protein	3.5	<b>-</b> 4.32
91	rc_AA486072_i_at		small inducible cytokine A5 (RANTES)	3.5	-3.88
92	rc_Z39983_s_at	Z39983	KIAA0561 protein	3.5	-5.56
93	rc_F02333_at	F02333	hypothetical protein FLJ20093	3.5	-2.23

Normal ve RPH	W/Symptoms Table	TARLE 3	49 1654540.1		
Up-regulated	Affy element		GenBank Name	Fold-change	t
94	rc_AA151210_at	AA151210	ESTs	3.5	-4.20
95	rc_N92239_at	N92239	Wnt inhibitory factor-1	3.5	-3.06
96	rc AA173223 at	AA173223	ESTs	3.5	-5.22
97	rc_T86148_s_at	T86148	pituitary tumor-transforming 1 interacting protein	3.5	-2.15
98	AA214688_at	AA214688	eukaryotic translation initiation factor 4B	3.5	-3.13
99	_	AA216589	ESTs	3.5	-4.40
100	rc_AA446661 at	AA446661	hypothetical protein FLJ10970	3.4	-3.69
101	AA082546 at	AA082546	ESTs	3.4	-4.12
102	rc_W46395 at	W46395	chromobox homolog 6	3.4	-2.41
103	rc_AA401433_at	AA401433	ESTs	3.4	-3.17
104	D62965_at	D62965	ESTs	3.4	-2.07
105	rc_AA057829 s at		growth arrest-specific 6	3.4	-2.00
106	rc_AA009755 at	AA009755	ESTs	3.3	-4.77
107	AA247204 at	AA247204	DEAD/H (Asp-Glu-Ala-Asp/His) box polypeptide 16	3.3	-2.85
108	D13628_at	D13628	angiopoietin 1	3.3	-4.86
109	rc_N59866 at	N59866	ESTs	3.3	-4.39
110	rc_AA406371_at	AA406371	ESTs	3.3	-4.98
111	rc N67876 s at	N67876	insulin-like growth factor 1 (somatomedin C)	3.3	-3.06
112	M84526 at	M84526	D component of complement (adipsin)	3.3	-3.06
113	rc_AA234095 at	AA234095	hypothetical protein FLJ20701	3.3	-3.78
114	rc_D60074_s_at	D60074	cadherin 10 (T2-cadherin)	3.3	-5.05
115	rc_T49602 s at	T49602	ESTs	3.3	-3.36
116	rc_n22006_s_at	n22006	ESTs	3.3	-3.88
117	rc_F04112_f_at	F04112	ESTs	3.3	-3.26
118	rc_T64223_s_at	T64223	carboxypeptidase A3 (mast cell)	3.3	-2.97
119	U23946_at	U23946	RNA binding motif protein 5	3.2	-3.48
120	rc_AA358038_at	AA358038	SH3-binding domain glutamic acid-rich protein like	3.2	-3.21
121	rc_AA019433 at	AA019433	ESTs	3.2	-3.88
122	X03689_s_at	X03689	eukaryotic translation elongation factor 1 alpha 1	3.2	-1.91
123	rc_H17550 at	H17550	ESTs	3.2	-2.90
124	rc_AA047880_at	AA047880	prothymosin, alpha (gene sequence 28)	3.2	-5.88
125	rc_AA084138_at	AA084138	ESTs	3.2	-7.93
126	rc_AA599365_at	AA599365	decorin	3.2	-4.42
127	rc_N91971_f_at	N91971	retinol-binding protein 1, cellular	3.2	-4.13
128	rc_T62873_at	T62873	ESTs	3.2	-2.12
129	rc_N49899_at	N49899	ESTs	3.2	-3.73
130	AA298981_at	AA298981	fibulin 5	3.2	-6.06
131	rc_AA479286_at	AA479286	ESTs	3.2	-3.54
132	J04111_at	J04111	v-jun avian sarcoma virus 17 oncogene homolog	3.2	-5.47
133	rc_AA465491_at	AA465491	Mad4 homolog	3.2	-2.75
134	W28548_at	W28548	ESTs	3.2	-3.59
135	AA308998_at	AA308998	endothelial differentiation-related factor 1	3.2	-2.89
136	rc_AA488432_at	AA488432	phosphoserine phosphatase	3.2	-3.48
137	rc_AA598991_at	AA598991	amyloid beta (A4) precursor protein-binding, family A, member 2 (X11-like)	3.1	-4.51
138	AA463311_at	AA463311	hypothetical protein similar to mouse Fbw5	3.1	-2.57
139	rc_AA147224_at	AA147224	ESTs	3.1	-4.41
140	rc_AA609504_at	AA609504	fibronectin leucine rich transmembrane protein 2	3.1	-3.81
141	U20734_s_at	U20734	jun B proto-oncogene	3.1	-3.37
142	U06863_at	U06863	follistatin-like 1	3.1	-2.48

50 Normal vs. BPH W/Symptoms Table TABLE 3 GenBank ID GenBank Name Up-regulated Affy element Fold-change t W51743 143 W51743 at 3.1 -2.95TIA1 cytotoxic granule-associated RNA-binding protein 144 rc\_AA465093 at AA465093 3.1 -5.34145 rc\_AA219100\_at AA219100 DKFZP586P2421 protein 3.1 -4.09**ESTs** 146 rc\_R42424\_at R42424 3.1 -3.82**ESTs** -2.23 147 rc\_W73038\_at W73038 3.1 148 AA091278 at AA091278 hypothetical protein FLJ10793 3.1 -2.75 PRO0518 protein 149 rc\_AA620289\_at AA620289 3.1 -2.55 prostate cancer associated protein 1 150 rc AA149579 at AA149579 3.1 -2.66small inducible cytokine A5 (RANTES) 151 M21121\_at M21121 3.1 -4.97**ESTs** 152 rc\_AA427890\_at AA427890 3.1 -4.32 153 M34516\_r\_at M34516 immunoglobulin lambda-like polypeptide 1 3.1 -3.47zinc finger protein 216 154 rc AA233347 at AA233347 3.1 -2.43155 W74533 latrophilin rc\_W74533\_at 3.1 -3.51 156 rc\_AA029597\_at AA029597 bone morphogenetic protein 7 (osteogenic protein 1) 3.1 -3.80 thymosin, beta, identified in neuroblastoma cells 157 rc\_N91887\_s\_at N91887 3.1 -4.47**ESTs** 158 rc\_AA205724\_at AA205724 3.0 -6.70 P311 protein 159 U30521\_at U30521 3.0 -6.06 protein kinase C, beta 1 160 X07109 at X07109 3.0 -4.90 potassium voltage-gated channel, KQT-like subfamily, 161 D82346\_at D82346 3.0 -3.49member 2 162 rc\_AA478962\_at AA478962 **ESTs** -3.35 3.0 matrix metalloproteinase 23A,matrix metalloproteinase 163 rc\_AA151428\_s\_at AA151428 3.0 -2.78 23B 164 rc\_AA130349\_at AA130349 **ESTs** 3.0 -2.01 granzyme A (granzyme 1, cytotoxic T-lymphocyte-165 M18737\_rna1\_at M18737 3.0 -5.90 associated serine esterase 3) **ESTs** 166 rc N91461 at N91461 3.0 -3.43 **ESTs** 167 rc\_AA045481\_at AA045481 3.0 -3.70U91903\_at frizzled-related protein 168 U91903 3.0 -4.73stromal cell-derived factor 1 169 U19495 s at U19495 3.0 -4.38tryptase, alpha,tryptase, beta (tryptase II) 170 M33493\_s\_at M33493 3.0 -3.12progesterone binding protein 171 Y12711\_at Y12711 3.0 -2.33**ESTs** 172 rc\_N58172\_at N58172 3.0 -2.53173 M12529 at M12529 apolipoprotein E 3.0 -1.92**ESTs** 174 rc\_AA412505\_at AA412505 3.0 -3.35glycoprotein M6B 175 U45955 U45955 at 3.0 -4.09176 rc\_H56673\_at H56673 3.0 -4.25 177 L33799 at L33799 procollagen C-endopeptidase enhancer 3.0 -4.72 **ESTs** 178 rc\_Z40186\_at Z40186 3.0 -2.22 eukaryotic translation initiation factor 3, subunit 7 (zeta, 179 AA094800 at AA094800 2.9 -2.5666/67kD) minichromosome maintenance deficient (S. cerevisiae) 180 D21063 at D21063 2.9 -5.27 2 (mitotin) **ESTs** 181 AA412049 rc\_AA412049\_at 2.9 -2.63 182 rc\_AA599661\_at AA599661 2.9 -8.62 collagen, type VII, alpha 1 (epidermolysis bullosa, 183 L02870\_s\_at L02870 2.9 -4.69 dystrophic, dominant and recessive) rc\_AA232266\_s\_at AA232266 184 2.9 -3.22 glutathione S-transferase M5 185 L02321\_at L02321 2.9 -3.33 SEC14 (S. cerevisiae)-like 2 186 rc\_AA428325\_at AA428325 2.9 -3.52 f-box and leucine-rich repeat protein 5 187 D82534 at D82534 2.9 -2.20KIAA0657 protein 188 T32113 2.9 rc\_T32113\_at -2.47

51 Normal vs. BPH W/Symptoms Table TABLE 3 1654540.1 GenBank ID GenBank Name Up-regulated Affy element Fold-change t 189 R10896 cytochrome c oxidase subunit VIIa polypeptide 2 like rc\_R10896\_at 2.9 -1.99 rc\_AA019034\_i\_at AA019034 190 2.9 -4.40 **ESTs** 191 D28423\_at D28423 2.9 -2.31 **ESTs** AA609943 192 rc\_AA609943\_at 2.9 -3.86 **ESTs** 193 W69302\_at W69302 2.9 -2.68 194 H01824 GATA-binding protein 2 rc\_H01824\_f\_at 2.9 -3.82 **ESTs** 195 rc\_T67105\_s\_at T67105 2.9 -5.49 H1 histone family, member X 196 rc\_AA426372\_s\_at AA426372 2.9 -2.53 197 **ESTs** rc\_T98288\_f\_at T98288 2.9 -2.66 **ESTs** 198 rc\_N63047\_at N63047 2.9 -5.25 GCN5 (general control of amino-acid synthesis, yeast, U57316\_at 199 U57316 2.9 -3.59 homolog)-like 2 alpha-2-macroglobulin

2,9

-1.76

200

rc\_AA219304\_s\_at AA219304

		•	52		
Normal vs. BPH	W/Symptoms Table		1654540.1		
Down-regulated	Affy element	GenBank ID	GenBank Name	Fold-change	t
1	rc_T40895_at	T40895	protein tyrosine phosphatase type IVA, member 1	16.5	5.19
2	rc_N80129_i_at	N80129	metallothionein 1L	12.6	3.54
3	rc_AA460914_at	AA460914	ESTs	7.4	4.58
4	rc_AA234996_s_at	AA234996	cytochrome c oxidase subunit VIa polypeptide 2	7.2	4.10
5	X66141_at	X66141	myosin, light polypeptide 2, regulatory, cardiac, slow	6.6	3.80
6	AA234634_f_at	AA234634	CCAAT/enhancer binding protein (C/EBP), delta	6.2	4.35
7	rc_AA419011_at	AA419011	prostate androgen-regulated transcript 1	6.1	3.87
8	rc_N94303_at	N94303	ESTs	5.8	5.96
9	M20543_at	M20543	actin, alpha 1, skeletal muscle	5.5	3.20
10	rc_AA085943_s_at	AA085943	troponin T1, skeletal, slow	5.5	3.02
11	X06825_at	X06825	tropomyosin 2 (beta)	5.2	3.35
12	AB000584_at	AB000584	prostate differentiation factor	5.1	3.80
13	M19309_s_at	M19309	troponin T1, skeletal, slow	5.0	3.41
14	rc_AA040433_at	AA040433	DKFZP586N2124 protein	5.0	2.62
15	rc_N32748_at	N32748	ESTs	5.0	3.36
16	rc_AA227926_at	AA227926	ESTs	4.8	5.39
17	rc_AA457566_at	AA457566	ESTs	4.7	4.22
18	rc_AA026641_s_at	AA026641	secretory leukocyte protease inhibitor (antileukoproteinase)	4.6	2.09
19	rc_AA053424_at	AA053424	serine/threonine protein kinase MASK	4.5	4.16
20	V00594_at	V00594	metallothionein 2A	4.5	3.71
21	rc_R16983_at	R16983	ESTs	4.5	3.23
22	U75272_s_at	U75272	progastricsin (pepsinogen C)	4.4	4.57
23	rc_T94447_s_at	T94447	cortic al thymocyte receptor (X. laevis CTX) like	4.4	3.50
24	U08021_at	U08021	nicotinamide N-methyltransferase	4.4	2.41
25	J03910_ma1_at	J03910	metallothionein 1G	4.3	2.79
26	rc_AA236545_at	AA236545	ESTs	4.2	2.41
27	rc_AA211443_at	AA211443	ESTs	4.2	4.49
28	rc_AA398908_at	AA398908	ESTs	4.2	2.64
29	X57129_at	X57129	H1 histone family, member 2	4.2	3.88
30	M21665_s_at	M21665	myosin, heavy polypeptide 7, cardiac muscle, beta	4.1	3.61
31	X65614_at	X65614	S100 calcium-binding protein P	4.1	4.03
32	rc_AA197112_r_at	AA197112	putative nuclear protein	4.1	3.07
	M99487_at	M99487	folate hydrolase (prostate-specific membrane antigen) 1	4.0	2.65
34	X04201_at	X04201	neurotrophic tyrosine kinase, receptor, type 1	3.9	2.87
35	X05451_s_at	X05451	ESTs	3.9	3.26
36	rc_AA435720_i_at	AA435720	tubulin, alpha 2	3.9	2.20
37	rc_N92502_s_at	N92502	ESTs	3.8	3.11
	L77701_at	L77701	COX17 (yeast) homolog, cytochrome c oxidase assembly protein	3.8	3.97
_	HG2157-HT2227_at			3.8	4.08
	X76717_at	X76717	metallothionein 1L	3.8	5.82
41	HG1067-HT1067_r_a	HG1067-HT106		3.7	3.02
42	rc_AA599331_at	AA599331	CGI-119 protein, uncharacterized bone marrow protein BM039	3.6	4.90
		M20642	ESTs	3.6	3.48
	rc_AA055163_at	AA055163	calsequestrin 2, cardiac muscle	3.6	3.66
	_	AA127946	DKFZP586B2022 protein	3.6	4.40
	_	AA022886	retinal degeneration B beta	3.6	3.51
47	rc_AA342337_at	AA342337	ESTs	3.5	2.57

			55			
	W/Symptoms Table			54540.1		
Down-regulated		GenBank ID	GenBank Name		Fold-change	t
48	X02544_at	X02544	orosomucoid 1		3.5	1.92
49	rc_T73433_s_at	T73433	angiotensinogen		3.5	3.10
50	M21494_at	M21494	creatine kinase, muscle		3.4	2.46
51	rc_AA488072_s_at	AA488072	cardiac ankyrin repeat protein		3.4	2.78
52	rc_AA293187_s_at	AA293187	B-cell CLL/lymphoma 3		3.4	1.62
53	rc_AA599522_r_at	AA599522	squamous cell carcinoma antigen recognised cells	by T	3.4	3.03
54	rc_AA405488_at	AA405488	ESTs		3.4	2.57
55	rc_AA461453_at	AA461453	calcium blnding protein Cab45 precursor,		3.4	3.10
56	rc_AA609006_at	AA609006	ESTs		3.4	2.30
57	rc_N24761_at	N24761	TU12B1-TY protein		3.4	3.89
58	rc_AA432162_at	AA432162	DKFZP586B2022 protein		3.4	2.78
59	X06256_at	X06256	integrin, alpha 5 (fibronectin receptor, alpha polypeptide)		3.4	4.51
60	rc_AA045825_at	AA045825	ESTs		3.3	3.90
61	rc_AA478778_at	AA478778	ESTs		3.3	4.37
62	rc_N80129_f_at	N80129	metallothionein 1L		3.2	3.60
63	rc_AA182030_at	AA182030	pyruvate dehydrogenase kinase, isoenzyme	4	3.2	3.72
64	rc_AA102489_at	AA102489	hypothetical protein FLJ10337		3.2	2.20
65	rc_R46074_at	R46074	transforming, acidic coiled-coil containing prof	tein 2	3.2	3.38
66	rc_AA599522_f_at	AA599522	squamous cell carcinoma antigen recognised cells	by T	3.2	2.36
67	rc_AA165313_at	AA165313	ESTs		3.2	2.76
68	rc_AA429636_at	AA429636	hexokinase 2		3.2	3.12
69	rc_R71792_s_at	R71792	thrombospondin 1		3.1	2.31
70	U05861_at	U05861	aldo-keto reductase family 1, member C1 (dihydrodiol dehydrogenase 1; 20-alpha (3-alphydroxysteroid dehydrogenase), aldo-keto red family 1, member C2 (dihydrodiol dehydrogen bile acid binding protein; 3-alpha hydroxysterodehydrogenase, type III)	uctase ase 2;	3.1	2.62
71	rc_AA410311_at	AA410311	ESTs		3.1	3.52
72	rc_AA505136_at	AA505136	ESTs		3.1	3.00
73	rc_T68873_f_at	T68873	metallothionein 1L		3.0	3.18
74	X00371_rna1_at	X00371	myoglobin		3.0	2.18
75	rc_AA099820_at	AA099820	ESTs		3.0	3.08
	rc_T90190_s_at	T90190	H1 histone family, member 2		3.0	3.48
77	rc_AA227936_f_at	AA227936	parathymosin		3.0	1.76
78	X90568_at	X90568	titin		3.0	2.83
79	rc_AA004699_at	AA004699	orphan G-protein coupled receptor		3.0	2.23
80	rc_F03969_at	F03969	ESTs		2.9	2.53
81	X93036_at	X93036	FXYD domain-containing ion transport regulat	tor 3	2.9	2.91
82	rc_R91484_at	R91484	ESTs		2.9	6.43
83	rc_AA025370_at	AA025370	KIAA0872 protein		2.9	2.87
84	X51441_s_at	X51441	serum amyloid A1		2.9	1.78
85	X64177_f_at	X64177	metallothionein 1H		2.9	3.36
86	rc_AA255480_at	AA255480	ECSIT		2.9	2.38
87	rc_AA476944_at	AA476944	ESTs		2.8	4.26
88	U78294_at	U78294	arachidonate 15-lipoxygenase, second type		2.8	1.82
89	rc_AA045487_at	AA045487	ESTs		2.8	2.75
90	rc_N74291_at	N74291	ESTs		2.8	1.88
91	rc_N91973_at	N91973	hypothetical protein,three prime repair exonuc	lease 1	2.8	1.97

			54		
Normal vs. BPH	W/Symptoms Table	TABLE 4	1654540.1		
Down-regulated	Affy element	GenBank ID	GenBank Name	Fold-change	t
92	D81655_at	D81655	ESTs	2.8	1.89
93	U53225_at	U53225	sorting nexin 1	2.8	3.16
94	rc_H77597_f_at	H77597	metallothionein 1H	2.8	2.98
95	K02215_at	K02215	angiotensinogen	2.8	3.05
96	rc_AA464728_s_at	AA464728	ESTs	2.7	3.80
97	rc_W49708_at	W49708	ESTs	2.7	3.52
98	rc_AA453435_at	AA453435	ESTs	2.7	4.78
99	rc_D11824_at	D11824	ESTs	2.7	3.70
100	rc_T56281_f_at	T56281	RNA helicase-related protein	2.7	2.62
101	rc_AA182882_at	AA182882	titin-cap (telethonin)	2.7	1.85
102	rc_AA447522_at	AA447522	ESTs	2.7	3.27
103	rc_N26904_at	N26904	FK506 binding protein precursor	2.7	3.21
104	rc_AA131919_at	AA131919	putative type II membrane protein	2.7	4.15
105	rc_R89840_at	R89840	ESTs	2.7	2.23
106	rc_W31470_at	W31470	thyroid hormone receptor-associated protein, 95-kD subunit	2.7	2.85
107	rc_W92207_at	W92207	ESTs	2.7	4.07
108	U96094_at	U96094	sarcolipin	2.7	2.23
109	rc_W70131_at	W70131	ESTs	2.7	3.64
110	rc_AA435720_f_at	AA435720	tubulin, alpha 2	2.7	1.98
111	rc_AA284879_at	AA284879	ESTs	2.7	1.74
112	rc_H22453_at	H22453	ESTs	2.7	4.20
113	D14826_s_at	D14826	cAMP responsive element modulator	2.6	4.13
114	rc_N93798 at	N93798	protein tyrosine phosphatase type IVA, member 3	2.6	3.12
115	U41804 at	U41804	putative T1/ST2 receptor binding protein	2.6	4.37
116	rc_W20486_f at	W20486	chromosome 21 open reading frame 56	2.6	2.74
117	rc_AA055768 at	AA055768	CGI-119 protein	2.6	2.13
118	rc_AA447977 s at	AA447977	ESTs	2.6	3.22
119	AA380393_at	AA380393	SEC7 homolog	2.6	2.29
120	rc_N29568_at	N29568	thyroid hormone receptor-associated protein, 150 kDa subunit	2.6	2.46
121	rc_AA426374_f_at	AA426374	tubulin, alpha 2	2.6	3.20
122	rc_H94471_at	H94471	occludin	2,6	2.19
123	rc_AA252219_at	AA252219	ESTs	2.6	3.83
124	rc AA402000 at	AA402000	ESTs.	2.6	2.29
125	rc Z38744 at	Z38744	putative gene product	2.6	4.18
126	AA045870 at	AA045870	ESTs	2.6	2.26
127	rc_R38678_at	R38678	ESTs	2.6	4.16
128	R39467_f_at	R39467	NEU1 protein	2.6	2.79
129	AA455001_s_at	AA455001	CGI-43 protein	2.6	5.34
130	rc_AA292328_at	AA292328	activating transcription factor 5	2.6	2.88
131	X57348_s_at	X57348	stratifin	2.6	2.48
132	rc_T95005_s_at	T95005	ESTs	2.5	
133	AA410355 at	AA410355	ribosomal protein S6 kinase, 70kD, polypeptide 2	2.5 2.5	3.30 2.31
134	AA036900_at	AA036900	ESTs	2.5 2.5	
135	rc_F02204_at	F02204	BAI1-associated protein 2		2.45
136	U26173_s_at	U26173	nuclear factor, interleukin 3 regulated	2.5	2.26
137	rc_AA477767_at	AA477767	ESTs	2.5	3.91
138	rc_AA504805_s_at	AA504805	interferon stimulated gene (20kD)	2.5	3.17
139	rc_R33627_i_at	R33627	ESTs	2.5	3.79
100	1.00021_i_dL	100021		2.5	1.99

Normalus DDU	W/Symptoms Table	TARLE 1	33	1654540.1		
Down-regulated		GenBank ID	GenBank Name		Fold-change	t
140	rc_T40995_f_at		alcohol dehydrogenase 3 (class I), gamma	ı	2.5	2.15
141	rc_R00144_at	R00144	polypeptide ESTs		2.5	2.69
142	U02020_at	U02020	pre-B-cell colony-enhancing factor		2.5	4.20
143	rc_AA287832_at	AA287832	ESTs		2.5	3.80
144	AA429539 f at	AA429539	hypothetical protein		2.5	2.35
145	rc_H05084 at	H05084	GDP-mannose pyrophosphorylase B		2.5	2.23
146	rc_AA405616_at	AA405616	ESTs		2.5	3.33
147	AA455381_at	AA455381	aldehyde dehydrogenase 5 family, member (succinate-semialdehyde dehydrogenase)		2.4	2.60
148	M13955_at	M13955	keratin 7		2.4	2.22
149	rc_AA180314_at	AA180314	ESTs		2.4	2.53
150	M37984_rna1_at	M37984	troponin C, slow		2.4	2.10
151	M61764_at	M61764	tubulin, gamma 1		2.4	3.48
152	rc_AA150920_at	AA150920	KIAA0539 gene product		2.4	4.11
153	X65965_s_at	X65965	superoxide dismutase 2, mitochondrial		2.4	2.37
154	X93510_at	X93510	LIM domain protein		2.4	2.39
155	rc_N48056_s_at	N48056	folate hydrolase (prostate-specific membra antigen) 1	ane	2.4	1.80
156	rc_N26713_s_at	N26713	ESTs		2.4	3.87
157	rc_AA282247_at	AA282247	ESTs		2.4	3.17
158	rc_D80617_at	D80617	KIAA0596 protein		2.4	2.02
159	rc_F02245_at	F02245	monoamine oxidase A		2.4	2.79
160	rc_R58878_at	R58878	ESTs		2.4	2.80
161	rc_W45531_at	W45531	ESTs		2.4	4.17
162	L25270_at	L25270	SMC (mouse) homolog, X chromosome		2.4	3.26
163	rc_W88568_at	W88568	glycogenin 2		2.4	1.90
164	rc_AA070752_s_at	AA070752	insulin receptor substrate 1		2.4	2.87
165	U24169_at	U24169	JTV1 gene,hypothetical protein PRO0992		2.4	3.41
166	rc_T15423_s_at	T15423	2',3'-cyclic nucleotide 3' phosphodiesteras	6 <b>e</b>	2.4	1.71
167	X78706_at	X78706	carnitine acetyltransferase		2.4	3.51
168	rc_T10695_i_at	T10695	enigma (LIM domain protein)		2.4	1.52
169	rc_AA430388_at	AA430388	HSPC160 protein		2.4	5.04
170	M68519_rna1_at	M68519	surfactant, pulmonary-associated protein		2.4	3.89
171	rc_AA421562_at	AA421562	anterior gradient 2 (Xenepus laevis) homo	olog	2.4	1.80
172	rc_T97243_at	T97243	prenyl protein protease RCE1		2.4	2.46
173	rc_T15409_f_at	T15409	ESTs		2.3	3.76
174	rc_T62918_at	T62918	ESTs		2.3	2.59
175	rc_R15108_at	R15108	ESTs		2.3	2.74
176	AA454908_s_at	AA454908	KIAA0144 gene product		2.3	2.77
177	rc_N64683_at	N64683	CGI-119 protein		2.3	2.27
178	rc_H99035_at	H99035	ESTs	۵۱	2.3	4.34
179	Y08374_rna1_at	Y08374	chitinase 3-like 1 (cartilage glycoprotein-3	9)	2.3	2.94
180	rc_AA236241_at	AA236241	ESTS		2.3	1.57
181	U52969_at	U52969	Purkinje cell protein 4		2.3	3.49
182	rc_R11526_f_at	R11526	parathymosin		2.3	1.71
183	rc_T15850_f_at	T15850	ESTS	nha	2.3	2.42
184	HG2259-HT2348_s_a		tubulin, alpha 1 (testis specific),tubulin, al ubiquitous ortholog of rat pippin	pıla,	2.3	2.91 1.45
185	rc_H15143_s_at	H15143	ESTs		2.3	
186	rc_AA101767_at	AA101767	2010		2.3	3.52

			00	-	
Normal vs. BPH	W/Symptoms Table	TABLE 4	1654540. <sup>-</sup>	I	
Down-regulated	Affy element	GenBank ID	GenBank Name	Fold-change	t
187	rc_AA193197_at	AA193197	sarcomeric muscle protein	2.3	1.98
188	U03688_at	U03688	cytochrome P450, subfamily I (dioxin-inducible), polypeptide 1 (glaucoma 3, primary infantile)	2.3	2.97
189	rc_R37774_at	R37774	cytochrome P450 retinoid metabolizing protein	2.3	4.11
190	rc_H81413_f_at	H81413	high-mobility group (nonhistone chromosomal) protein isoforms I and Y	2.3	3.12
191	X16354_at	X16354	carcinoembryonic antigen-related cell adhesion molecule 1 (biliary glycoprotein)	2.3	2.54
192	rc_AA457235_at	AA457235	ESTs	2.3	2.25
193	D13643_at	D13643	KIAA0018 gene product	2.3	1.78
194	rc_N30856_at	N30856	solute carrier family 19 (thiamine transporter), member 2	2.3	3.45
195	M26311_s_at	M26311	S100 calcium-binding protein A9 (calgranulin B)	2.3	2.37
196	rc_Z40556_at	Z40556	CGI-96 protein	2.3	2.39
197	rc_N79070_at	N79070	ESTs	2.3	1.43
198	Z69881_at	Z69881	ATPase, Ca++ transporting, ubiquitous	2.3	3.87
199	rc_D60755_s_at	D60755	ESTs	2.3	2.30
200	rc_N94424_at	N94424	retinoic acid receptor responder (tazarotene induced)	2.2	1.09

Table 5 57

1654523.1

	lated genes		egulated genes
Cluster	Fragment Name		Fragment Name
1	rc_AA256268_at	1	rc_AA227926_at
1	rc_AA188981_at	1	rc_AA398908_at
1	rc_AA173223_at	1	L77701_at
1	rc_AA216589_at	11	rc_AA599331_at
1	rc_AA234095_at	1	AA455001_s_at
1	rc_H17550_at	3	rc_AA022886_at
1	AA308998_at	3	rc_N24761_at
1	rc_AA488432_at	3	X06256_at
1	rc_AA427890_at	4	HG1067-HT1067_r_at
$\overline{1}$	rc_N91887_s_at	4	rc_AA127946_at
11	rc_AA045481_at	4	rc_AA405488_at
3_	rc_T23622_at	5	AA234634_f_at
3_	rc_T23490_s_at	5	X65614_at
3	rc_AA620289_at	5	rc_T73433_s_at
4	rc_H05704_r_at	5	rc_R91484_at
4	rc_AA436616_at	5	rc_N93798_at
4	rc_AA456147_at	6	rc_N94303_at
4	rc_f09748_s_at, AA495865_at	6	AB000584_at
4	rc_AA598982_s_at	6	rc_AA410311_at
4_	HG3543-HT3739_at	6	rc_F02245_at
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			rc_T68873_f_at
5	U62015_at	7	rc_N32748_at
5	rc_F13763_at	7_	V00594_at
5	rc_AA205724_at	7	J03910_rna1_at
5	U30521_at	7	X57129_at, rc_T90190_s_at
6	X52541_at	7	rc_AA182030_at
6	rc_AA281345_f_at, M62831_at	7	rc_AA505136_at
7	rc_n22006_s_at	7	X64177_f_at, rc_H77597_f_at
7	rc_R42424_at	7	rc_AA101767_at

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1654537.1

Table 6

	Prostatic tissues	Cell Line				
		BRF-55T	PZ-HPV7	BPH-1	LNCaP	
Up-regulated genes	61	33	22	20	20	
Down-regulated genes	43	31	28	30	33	
Total	104	64	50	50	53	

WE CLAIM:

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- 1. A method of screening for or identifying an agent that modulates the onset or progression of benign prostatic hyperplasia (BPH), comprising:
- (a) preparing a first gene expression profile of BPH cells or BPH-like cell population;
  - (b) exposing the cells to the agent
  - (c) preparing second gene expression profile of the agent exposed cells;
    - (d) comparing the first and second gene expression profiles.

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and

- 2. A method of claim 1, wherein the gene expression profile comprises the expression levels for one or more genes that are differentially regulated in BPH cells compared to normal prostate cells.
- 3. A method of claim 1, wherein the agent modulates the expression levels for one or more genes in the BPH cells to levels close or similar to the expression level found in normal prostate cells.
- 4. A method of claim 1, wherein the gene expression profile comprises the expression levels in BPH cells for one or more genes in Tables 1-5.
  - 5. A method of claim 1, wherein the gene expression profile comprises the expression levels in BPH cells for one or more genes in Table 5.
- 6. A method of any one of claims 1-5, wherein the BPH cell is selected from the group consisting of prostate cells from a BPH patient, a cell line in Table 6 and a derivative thereof.

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- 7. A method of any one of claims 2-5, wherein the expression levels are for two or more genes.
- 8. A method of diagnosing the onset or progression of benign prostatic hyperplasia (BPH) in a subject comprising:
  - (a) detecting the expression levels of one or more genes in prostate cells from the subject that are differentially regulated compared to normal prostate cells.
- 9. A method of claim 8, wherein the expression levels are for one or more genes in Tables 1-5.
  - 10. A method of claim 8, wherein the expression levels are for two or more genes in Tables 1-5.
- 15 11. A method of claim 8, wherein the expression levels are for one or more genes in Table 5.
  - 12. A method of claim 8, wherein the expression levels are for two or more genes in Table 5.

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- 13. A method of differentiating benign prostatic hyperplasia (BPH) from prostate cancer in a subject comprising:
- (a) detecting the expression levels of one or more genes in prostate cells from the subject that are indicative of BPH rather than prostate cancer.

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14. A method of claim 13, wherein the expression levels are for one or more genes in Tables 1-5.

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- 15. A method of claim 13, wherein the expression levels are for two or more genes in Tables 1-5.
- 16. A method of claim 13, wherein the expression levels are for one or more genes in Table 5.
  - 17. A method of claim 13, wherein the expression levels are for two or more genes in Table 5.
- 18. A set of oligonucleotide probes, wherein each of the probes specifically hybridizes to a gene in Tables 1-5.
  - 19. A set of oligonucleotide probes, wherein each of the probes specifically hybridizes to a gene in Table 5.

20. A set of oligonucleotide probes of claim 18, wherein the set specifically hybridizes to nearly all the genes in Tables 1-5.

- 21. A set of oligonucleotide probes of claim 18, wherein the set specifically hybridizes to nearly all the genes in Table 5.
  - 22. A set of oligonucleotide probes of any one of claims 18-21, wherein the probes are attached to a solid support.
- 23. A set of oligonucleotide probes of claim 22, wherein the solid support is selected from the group consisting of a membrane, a glass support and a silicon support.

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- 24. A solid support onto which two or more oligonucleotide probes have been attached, wherein each of the probes specifically hybridizes to a gene in Tables 1-5.
- 25. A solid support of claim 24, wherein the probes specifically hybridize to nearly all of the genes in Tables 1-5
  - 26. A solid support onto which two or more oligonucleotide probes have been attached, wherein the probes specifically hybridize to a gene in Table 5.
- 10 27. A solid support of claim 26, wherein the probes specifically hybridize to nearly all of the genes in Table 5.
- 28. A solid support of any one of claims 24-27, wherein the solid support is an array comprising at least 10 different oligonucleotides in discrete locations per square centimeter.
  - 29. A solid support of claim 28, wherein the array comprises at least 100 different oligonucleotides in discrete locations per square centimeter.
- 30. A solid support of claim 28, wherein the array comprises at least 1000 different oligonucleotides in discrete locations per square centimeter.
  - 31. A solid support of claim 28, wherein the array comprises at least 10,000 different oligonucleotides in discrete locations per square centimeter.
    - 32. A computer system comprising:

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- a database containing information identifying the expression level in (a) benign prostatic hyperplasia (BPH) tissue or cells of a set of genes comprising at least two genes in Tables 1-5; and
  - (b) a user interface to view the information.

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- A computer system of claim 32, wherein the set of genes comprises at least 33. two genes in Table 5.
- A computer system of claim 32, wherein the database further comprises 34. 10 sequence information for the genes.
  - A computer system of claim 32, wherein the database further comprises 35. information identifying the expression level for the set of genes in normal prostate tissue or cells.

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- A computer system of claim 32, wherein the database further comprises 36. information identifying the expression level of the set of genes in prostate cancer tissue or cells.
- A computer system of claim 32, further comprising records including 37. 20 descriptive information from an external database, which information correlates said genes to records in the external database.
  - 38. A computer system of claim 37, wherein the external database is GenBank.

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A method of using a computer system of claim 32 to present information 39. identifying the expression levels in a tissue or cells of at least one gene in Tables 1-5, comprising the step of:

- (a) comparing the expression level of at least one gene in Tables 1-5 in the tissue or cells to the level of expression of the gene in the database.
- 40. A method of claim 39, wherein the expression levels of at least two genes are compared.
  - 41. A method of claim 39, wherein the expression levels of at least five genes are compared.
- 42. A method of claim 39, wherein the expression levels of at least ten genes are compared.
- 43. A method of claim 39, further comprising the step of displaying the expression levels of at lest one gene in the tissue or cell sample compared to the expression level in BPH.
  - 44. A method of monitoring the treatment of a patient with benign prostatic hyperplasia (BPH), comprising:
    - (a) administering a pharmaceutical composition to the patient;
  - (b) preparing a gene expression profile from a cell or tissue sample from the patient; and

- (c) comparing the patient gene expression profile to a gene expression profile from a normal prostate cells, or a BPH tissue or cell sample without treatment.
- 45. A method of claim 44, wherein the gene expression profile comprises the expression levels for one or more genes in Tables 1-5.

- 46. A method of claim 44, wherein the gene expression profile comprises the expression levels for one or more genes in Table 5.
- 47. A method of claim 45 or 46, wherein the expression levels are for two or more 5 genes.
  - 48. A method of any one of claims 1, 8, 12, 38 or 43, wherein the gene expression profile or gene expression level is detected by branched DNA (bDNA) method.
- 10 49. A computer readable storage medium storing a computer program for implementing an algorithm executing method of analyzing gene expression results; said method comprising:

respectively.

50.

15

- (a) converting the mean expression value for each gene to 0; and
- (b) converting the high and low expression values to 1 and -1,

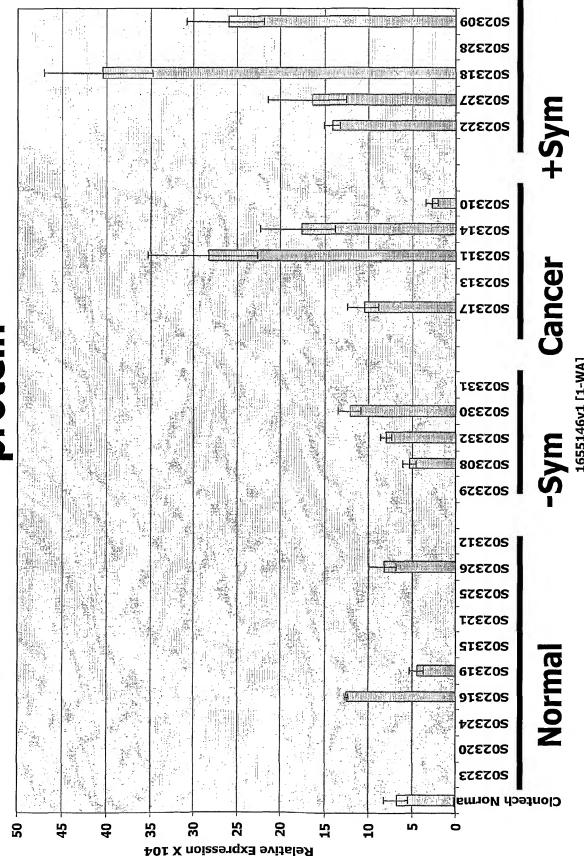
The medium of claim 49, wherein the method further comprises the step of:

- (c) clustering the converted expression values to identify sets of genes with similar expression patterns.

1/6

## H Figure 1: N91971, cellular retinol binding elos nu legeles s DUELO RIBUILIEN SNOTO sheet. Oun> UBB/US Snusyl NOLIEN SIOS Soello's protein ounsetul lieus Phos Bully EJURORIA URIE REPORT Unillederes DIONAL SI COLO (01/> Puelo Alenges SEROLEY. Puelo levelor 3000 2000 1500 0 2500 1000 500 Relative Expression X10 $^{4}$





elognin leteles so

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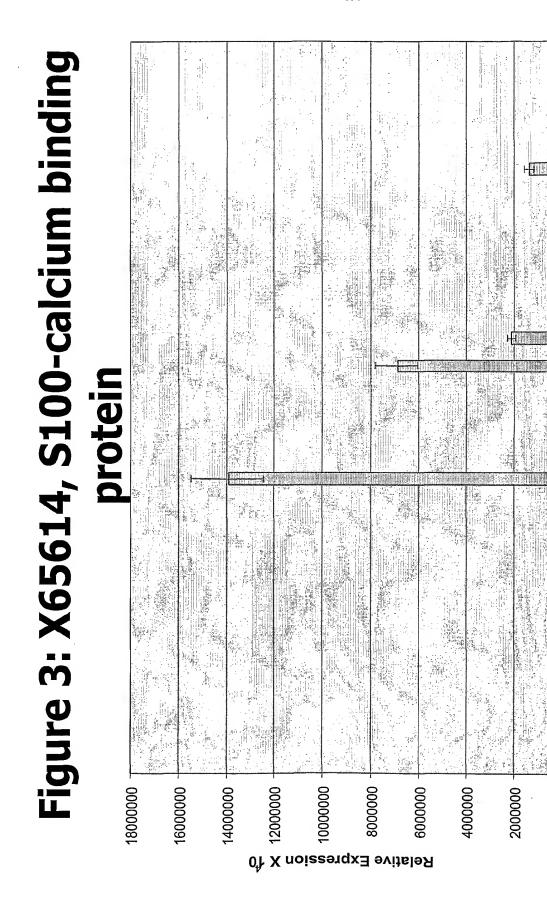
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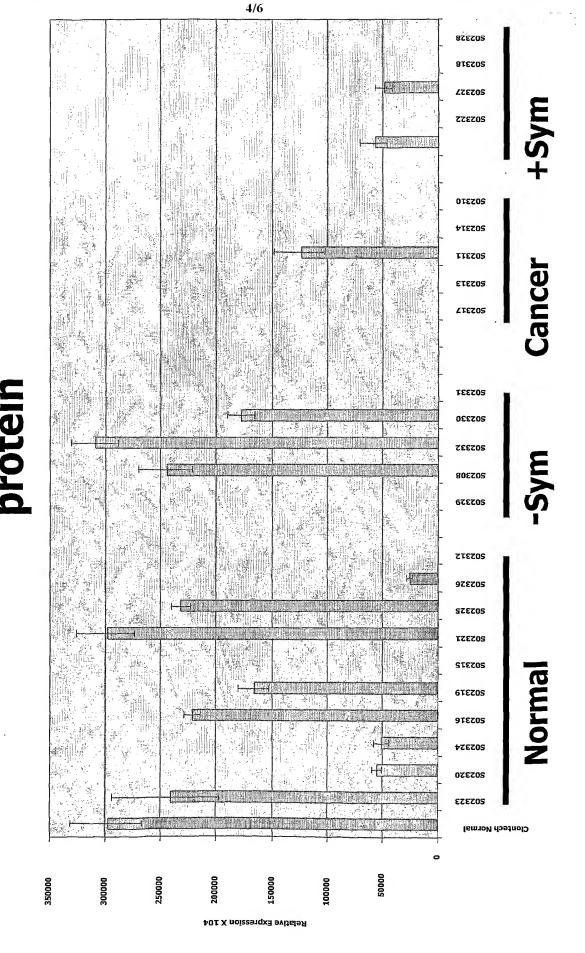
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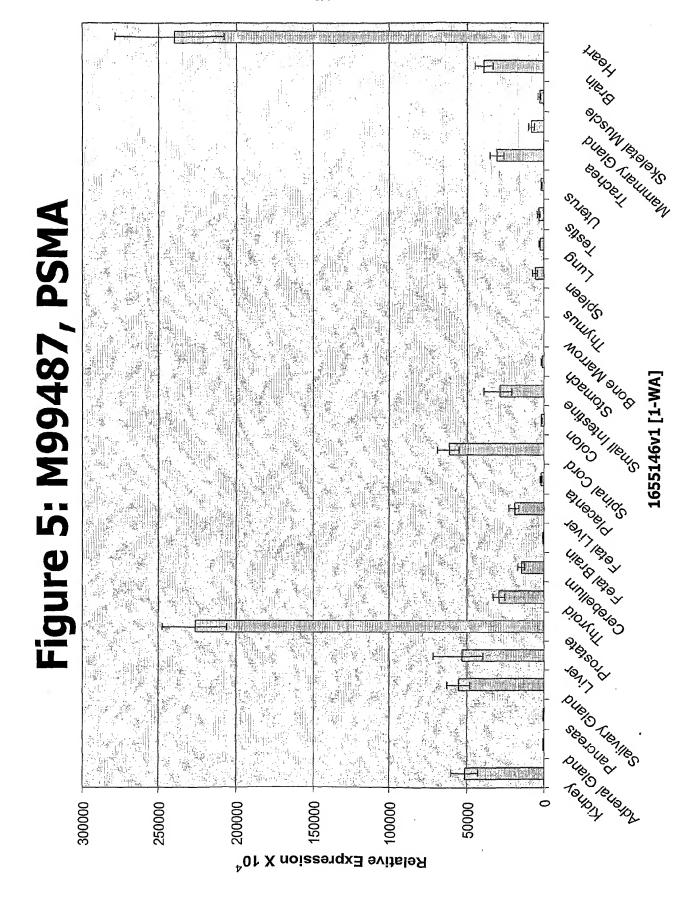
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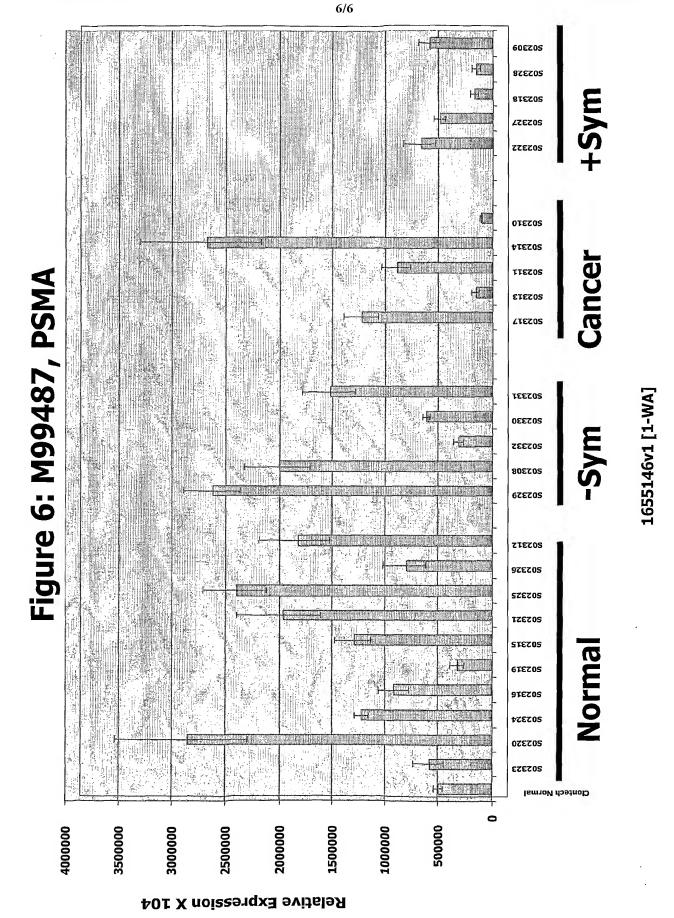


1655146v1 [1-WA]

## Figure 4: X65614, S100-calcium binding







## SEQUENCE LISTING

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<110> Munger, William E.
      Kulkarni, Prakash
      Getzenberg, Robert H.
      Waqa, Iwao
      Yamamoto, Jun
<120> Identifying Drugs for and Diagnosis of Benign Prostatic
      Hyperplasia Using Gene Expression Profiles
<130> 44921-5029-WO
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ctttgctacc ttctgcttgt tgagttgttt tggcattcat attaaaagcc agcatctcac 60
tatttattga caggttgggc tgtgtgtgtg cgcatgtgtg tatacatttc caggcgtgcc 120
tgtgtcctgt agctttttaa aaggaaaccc agtcatccca ctatgaatct ggcatcttct 180
tatgcttcta gtgttttggc canaca
<210> 15
<211> 494
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA039935
<220>
<221> unsure
<222> (1)..(494)
\langle 223 \rangle n = a or c or g or t
<400> 15
ttcagttttt ggtcatttta attgtaaaaa ccaagacatt tatataaata agaccgctgt 60
gtaaaatacg attcaccctt ctacgaaaac ccttttccca cactcgaaan gaanatagaa 120
aacccagcag agagcagtac aantcagcat geggteeeng atagetgaag tetegggeng 180
gccagtggtt ccctgcggaa nagccttcgt nggtgganag nactcctggc ccaggtggnc 240
ccaccagann ntcnntgacc ntctcnanga gacttgcnag gtangcagct cccnnacacc 300
agccccttgn gtctcaantn tacgggtcca aggagggac gggaaaggct gcttggtccc 360
caccaagget tggggggetg ggggggeetg etggeecagt gaagatgeag tggtetgtte 420
agcctggggt caagttgggg gaaagggttt ctgaggggtc agcacctccc cagaggacaa 480
ggagagaagc tgcn
<210> 16
<211> 421
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA040433
<220>
<221> unsure
<222> (1)..(421)
<223> n = a or c or g or t
<400> 16
ttttgagggt aatgcaactt cttatttatt aatatataat aacaacaatt atacagctca 60
tatctgcaac tgttaggtct ttgttatgtc ttggtcactt tgtctqqact qqccgtqacc 120
ttcagctcca gggtctgggc taggaagacg ttccagtgac cttcgtgggg gccagcgagc 180
agteggaagt getgtgeete tttetggaag tettgettee tgaetttett gatetgagte 240
aagtggaaga ttctggctgt gtggccttgg cagggtactt cacctctctq aqcctcaqtt 300
tecteatett ttaccagett ecagaggtag atetecacea agtecqagge etengtgtte 360
ccaggggcaa agcgacgnag gttngtctng ggctttgggg gataccggat gttttggacg 420
                                                                   421
```

<210> 17

```
<211> 486
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA040731
<220>
<221> unsure
<222> (1)..(486)
<223> n = a or c or g or t
<400> 17
ttccaagaag aacattttc tgtttattct tagaatgtga atttttttt tcaactcagg 60
gccaagtaca aacttttgat ttttgaaatt ttttcaactc agggccaagt acaatctttt 120
gatttaaaaa tttttttca tgaacaaacc atcagtagtt attaaggagc ccaagaaata 180
ggagatgtga aagcaggatt totttgtgtt tootttgaat gttgttattt tgagtattat 240
cattatcagg tagaggaaga aaggtaggct gggaagtagg tccttatgat atcttgacta 300
tggatcccag atttacattt cacctngtca cagagcacac ataatttaag ataaacatqt 360
caagaatgac ataaaccaga ggtaaacacc aaggagcttt acatttggaa ccngaaaata 420
aaaattagaa aaattattac cccatattaa taaccaaaaa attacttaaa ctcctaggnc 480
cccngg
<210> 18
<211> 546
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA043349
cttttatgca gagtttgatt atgctttatt tttaaaaatc acattcttcc ccattcccag 60
ccaacgaaca acactattca ttctgaaata agaaatgaa agaattttga gaagtcacac 120
aacattgctg taaatttcat tttttttt tttactaata aaacagatgc ttctttctca 180
gagatgggtt ttcactttca acatgcgtca tagcatctga ttttctgagc catcttggga 240
aatggagtct ttcctaatgt cattgaatgt ggtcaaagct atctacaaag cagagacagt 300
aggetettgg tgaateagtt tgggaaatte acaattaage agteteaggg agtgaaatte 360
cggggtctga tgagactgtg gaaaccatgt ggtactgtag ggagagcaca ggtttggatg 420
ccagacaaat atctaaatct aaccctaatc cactgcttat aagcttagtg attgttgcac 480
aagttgttta gcttctctga gcttagatac ctcactgtaa aatgggaata atacctcttt 540
ttagtg
<210> 19
<211> 353
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA043777
<220>
<221> unsure
<222> (1)..(353)
\langle 223 \rangle n = a or c or q or t
```

```
<400> 19
gaagttataa aagcttgttt ttctttatta gaatactttt ttcaattctg atttgtcaca 60
atttagattc tttttctaag aataagcaga aatttacaaa atttaatttt tatttataca 120
ttcatccqtt caatacacat ttcaagaaag ctgtattqna ccccttnnaq tnqqtaaqtt 180
ccagggccaa agaaccaaaa taaatccaag gagagagacc aacaaatgta tatttataac 240
acagagtaat aaaacacaaa taaatgtgga gttatttaag catgtaaqat gqtacatgct 300
ctaccaaggt atgggggctt ctctaagaca caagatcaga ttaaaqtctt qaa
<210> 20
<211> 382
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA044219
<220>
<221> unsure
<222> (1)..(382)
\langle 223 \rangle n = a or c or g or t
<400> 20
ttgcggggaa tcaggtaggg gcctttattg gccagcacac atctacctcc tggcatctqt 60
cacaagcatt tgcaggagta ggcggccct tcctctccat gtccccatcc ccaacctgaq 120
atgcgggagg gcctgggggc tcagagggaa gaactgaggc aagaagcccc gqtqatccag 180
tcagaggatt gggcagcctg acctcggggt ggggagccag cactngacaa caaggaggga 240
ggggcacagg agggctcccc gaggtttggt ccgggagggg gaggaaaact gcccctgcn 300
ctgtcaatct ctgcaatgtg ccgagcccca gctccttgan tccctcagtg cctttgggcc 360
tggatgctca ganagcagtt ga
<210> 21
<211> 428
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA045481
<220>
<221> unsure
<222> (1)..(428)
\langle 223 \rangle n = a or c or q or t
<400> 21
tttttttcag taatacagat gtctatttta ttaaaaaagt tacaaacagg tggactgcag 60
ggtcgtctta caaaatgaca agaatgaaat ctattggaaa aattttactt ttacaaatct 120
ttataggtaa ttgttcaatg tttgtacttg ttatttgaga ttttaccttt cactgataaa 180
gttacagtac attagatcca tgataatagg ttacattatt ttatttgcag agccctactg 240
cagtgatttg aacaactcct aaatagatgc cataataaag acaagacata tattgcattt 300
aatattaatt tattatccta ataagcaaca tgcaatctat tgaggaagct aaaataactt 360
ttggtcccct ttcttaaaat gtgctggaga aaccaccctt aaaatcactt tcccccggat 420
tccngcga
                                                                   428
<210> 22
<211> 328
<212> DNA
```

```
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA045487
ggaaagcatt ttcaaacttt atttacaact gtcacagtga caaaaagtag tttggaaaaa 60
aaaaaaatgct agtttctccc tgagcctcaa aaaagaacag atagaagtta caggaggttc 120
atctcacaac aggcattttt actgaaatac taggaatttt ttcaatacaa tcagttagaa 180
atacacacaa attacttgaa aaaaaaaaaa agaggaggcc agataggagc tcagccactt 240
gtccaagagc agetgggtcc ccccagcagg ctccaccgct gagggtcctg acattagctg 300
tcagccctg gcctgctcag actggcaa
                                                                   328
<210> 23
<211> 402
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA045503
<400> 23
ctgtgagact gtcctttatt gtgtatacag gttccagcgt cagggctctc ccacggcccc 60
ctccccagtc ctccccaag ggcccagagt ggtgggagtg agaggccacc ctaaggcaca 120
ctgaccagag aggcatggag ggaggaggct gacttgccct ggggacccct gctaactgag 180
acccaccett cccctccacc etgettetgt atgtgggaga cgaaaccaag agtcactggg 240
ggcagcaggc atttcccagg gttaaggctg atggaaggtc cctatcccag atgggagatg 300
ggggetttte ctatgactec ceccatecee cagetggaag acgtgggag gggtgcatag 360
ccttagagag gtagaatgag gggaaatact cctcagtgcc ca
<210> 24
<211> 437
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA045825
<220>
<221> unsure
<222> (1)..(437)
\langle 223 \rangle n = a or c or g or t
<400> 24
cagtgtagac cgtctttatt ggcaggtgtt aagagtgcaa aatatcaaca aacccagggg 60
aatacgcaag ggggtgggag tatggctccc ctaccccatg tgagagccct gtaaccaagc 120
cagtggggtg ggaacgttga cttgactgtn gcaaattcag gctcagcacc ttccaaagaa 180
caagctccca ggcaggaggg ctccttgcaa cacaaggggg aaaggagtgg caccctggaa 240
gggcctgggc tgcgacccac cctgggctgc ttggctcctg tatactgccc acctcaaccc 300
ctcaagagga aggcttcaca gctgggggta tgtagttcag agaacccggg ctaaacccag 360
ccctccccaa acccaggtta tctgcctcgg gcctcagttt ccctcctccc agtgattacc 420
caagttgggc ccatcag
                                                                   437
<210> 25
<211> 397
<212> DNA
```

```
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA045870
<220>
<221> unsure
<222> (1)..(397)
<223> n = a or c or g or t
<400> 25
gtttagagtc taaaactaaa acctaatcat ttnqtcacaq tqtaaaaaca aatqqaaata 60
acagctcaaa tcttcaaaat attactatag cattatgttt aaaataatct acaacaaaaa 120
tgtaccattt tcaagcagta ctacattagg agccctttta tagaaaataa tttcttcttt 180
accecegtte cagtgtgaat ctagtattet gttaacattt gtgtggcatt tggagtttgt 240
catececatt qaaggaaga cetteteaga catgaagcaa gggaaacata etgaatagtt 300
ttacacaaat ttgatctggc ttccatttgn ccccctcatt tcccaaatgt ttaaantgta 360
ttnggatttg ggattctcaa atggtataag ttggcct
<210> 26
<211> 564
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA046426
<220>
<221> unsure
<222> (1)..(564)
\langle 223 \rangle n = a or c or g or t
<400> 26
tttttttttt tttcacttta tcatttactt tttattgtgt tgcttgaagt acctatgtaa 60
tgcaagtatg tactgtacta aaatacctat atttccaaat aacatatgtg gtgtagccca 120
cagtetetge agaageatea tgagtaacet gtgeetttae actttacaat cegttattgg 180
ttgctgttaa aagtatgata acagatgaag aaaaaaaac taagtatgaa tacacttttc 240
caaacacgca catacacagc ttacaatgga atcccaatgg aaataagtga caacatctga 300
tgtagaatct ataaaatgta gactctgcaa taaaaagcca aaggacgtaa aaatatattt 360
taactttaaa aataacttag ttacagtaat actttgcctg tgtcttacca acatgtagct 420
gacagtcaaa attttgcaat atagatataa tatataggga tatataagaa ctacaagaaa 480
atccccaaaa cccataaagt tcaaatgtga aacagaaaag tttaacctgg agattcgcta 540
tggtgancta gccatatttg gaag
<210> 27
<211> 560
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. AA046840
<220>
<221> unsure
<222> (1)..(560)
\langle 223 \rangle n = a or c or g or t
```

```
<400> 27
tacaaatact qtaaaaatta atataaaaaa qtgaqcatgc tcagtctttt cctcttatct 60
acaatacaaa gggtttgtct gaaaagtctg gttttttttc tttttacaaa tqtaccttag 120
ctqcatcaac aggagtaaga tgtagaaaaa gctaccatta caaaaataat ttaaqqqaaa 180
ataaacacgt ttagcttctc tcgcagttta gtggtggtaa gtccaggctg tagcttcttt 240
gegeteetat gteecaagaa actgeagegg geacceggeg getetggetg egeagggeag 300
ggegegetee geteegggee gtegggtetg aggtatgggt egttgetgag teteteeege 360
cccggccgcg cgttaccggc agtctgctgt cccggcggcc ggcagaaggg cgggctgggc 420
agctgcttga agaactgccg gagggccagg tcccgcgtga ntgctccacg cgctggtgca 480
gttctcgttt cagcgacagc tcacaacttt gtgcantcct ggttgcgccg cttggcttgt 540
ggggtttgcn acgggatgtt
                                                                   560
<210> 28
<211> 464
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. AA047151
<220>
<221> unsure
<222> (1)..(464)
<223> n = a or c or g or t
<400> 28
agaaaaacca ccatcgtgtc acgtcgacga tgccaaatta tgttagcgtg acaganaaca 60
ccgtggggga ggaaggcagc agctgaagaa aaaagctcaa atgatctagt cactttcgat 120
actgtacttc agatgcgaaa tggatattcn gagtggaaac ctgacaaagt gcqcctgctt 180
tgatgtgaac tggtatagac aatgaccagt ggctgggtca gtgggatgtc tctctgtgag 240
cacaaaggct tatcaaatga cactaaagat aagttcaaca accatcacat tggaagggag 300
aaaggccgaa catttcatgt ttggccgggc atgtgagtgc acaagatgga aaqaqcqatt 360
ggagcatcct ggtataatta cccccattgt gctcttaatg gaaatttcaa aggacgggag 420
tattctgttg gttggtgtcc aggtttgtgg cactgttcca agag
<210> 29
<211> 413
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA047880
<400> 29
tacagagaat ataaaaatac attcacttta ttttaqaaaa atqaagactc atagagtaag 60
cttatcacaa actggcctat taggagtcac agaattcaca ggaaacaatt tctgaagacc 120
aggtgcctgc tgccacctct ccaagcaggc cagagtccag tagagaatgc gattcaggaa 180
gatggctcct cagagggcag ggaggttagc tacggaggcc gctcacgtgg aaatgtccag 240
tgaaccaatg ccaaggaaga agataaaatt ctctggggct gaccacaaca gtgggggtgg 300
ataaagacaa accacttgcc tgtacttctc atcttctatt tgttcatttc actgctggaa 360
ggtgacctct tttcccctaa tcttctttca acccagagag tttaagtctt ctc
<210> 30
<211> 431
<212> DNA
```

```
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA053424
<220>
<221> unsure
<222> (1)..(431)
\langle 223 \rangle n = a or c or g or t
<400> 30
tttgagcttt cagatttgct tttattggta gggaaattcc agagtgggga gccacccagg 60
aggagacagg ggtgccgagg cttctgggag tctggaagct cccggatgga gaggcttaca 120
gccccagcct tccccagcag gagcacaggc aggggactgg ccaagtctgt cagctcagag 180
caggaccggc ttcagggcct gacttcggtc tcctcttgac ccgccccgga ggcttgtggt 240
gggctctgtg tttgcagctc tcctgaacag agctagatga gggtgggagg cccccgttgg 300
ctcacacagt ggatgctacc atctccggcc tcttggatgt ggagctctgt gccagagtca 360
acagteteca gggtgggccg gaagttgttg taggcgntet caaggeegaa atetgetett 420
cctcagattc t
                                                                   431
<210> 31
<211> 451
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA055163
<400> 31
tttttcaaaa tatgagttta atgacagaat tagttagcta gtattccaca aaaagtattg 60
ctctattttc aaaaaatttg cacagtgtct tacacatgtg ctaaaagatt gagaaaataa 120
attagaaaat tatactgcac acttaacact aaatctacca agcacaatqt aacttttaga 180
cagctcagaa ggcactttgg gattttttt tttttcagtg cctcagggat cagtatgaac 240
tccaattatt gttgccctgg ccaattgtgg gagtactgat aactggagag ttaattgact 300
gctggataaa gcaatcttta atctaaatgg ggaaggctca ctagcagcta cagaggaagg 360
gggtattcag atcccagctt aaggctagga agccagctga cccaatcaga gacatqaacc 420
catcagaaaa atgtaaaagt tttcatcttt c
<210> 32
<211> 354
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA055768
<400> 32
tttttttttt tctgttcaaa aaaggtttta tccaaaaaag ttaatcaaga caagcaacag 60
atactgcaaa gcattatata cagcaccata gtccaggggc caaagaaatc aggagggct 120
gggcagtaga ggaattccat atattaatga atgtgagatt aagtatagag tgaaqacatt 180
aacacacaat ttctaatttc tgttaggcag aatgctcccc taccctgatg ccacagcctt 240
tcacgtttcc taaaccctag taacctctga tctccatctg cctcatcaac acgtcaccac 300
cetttgetet tettecaatt tagteacatg ttgggetgaa tttattteca etce
<210> 33
<211> 610
```

12

```
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. AA056121
<220>
<221> unsure
<222> (1)..(610)
\langle 223 \rangle n = a or c or g or t
<400> 33
ctcccctcc ctgctccaag ccggagggtt cctgaggtga cagcgcctgc aactgaaatt 60
tcagcagcgg gagaagatgg acaagagaaa gctcgggcga cggccatctt catccgataa 120
gaaagatgtt aaatgcaaaa ccagaggatg tccatgttca atcaccactg tccaaattca 180
gaagctcaga acgctggact ctccctttgc agtgggaaag aagcctaagg aataaagtca 240
tetetetaga ecataaaaat aaaaaacata teegagggtg teetgttact teeaagteat 300
caccagaaag gcaactcaaa gttatgttga cgaatgtcct atggacggat ttaggacgaa 360
aattcagaaa gaccctacct agaaacgatg ctaatttatg tgatgccaac aaggtgcaat 420
cagactcatt gccttcgaca tctgttgaca gcctagagac atgtcaaaaa ttagaacctc 480
ttcgccaaag ccttaattta tctgaaagga tnccagagtt atattgacga atgtctggga 540
acgggttagg aagaaatcct aaggncccac ctgtactgag ggaattggtg ttcagcaant 600
gcatcaggga
                                                                   610
<210> 34
<211> 404
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. AA057195
<220>
<221> unsure
<222> (1)..(404)
<223> n = a or c or g or t
<400> 34
agaaaaacca agtgtcttta ttcctcgatc gtttagtatg gcggtgggcg gcgcgcgcgg 60
gggagcctgg agcccaggga atcgacctgg agggccagtn gngggancgg agggtgcgag 120
gntcggctcc tccgcagccg gccctggagg ggttcttggg ggatcgcgcc aggccaaaaq 180
tetgeatggg eggeeegag cetecetgag eeggegege eegggnttng ggagaggeen 240
ctctgnncgc ggtgccgntg cgggcccggg tgcggctc gcccaagggc taaggtgccc 300
cgtctcaggc gagaccccag gagcccgccg ccccgctgt ctcttcagcc gacgtagaca 360
cgtngggccg ggaaccccag tcttaacgcg tgttcaagct ctgg
                                                                   404
<210> 35
<211> 491
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. AA057829
<220>
<221> unsure
```

```
<222> (1)..(491)
\langle 223 \rangle n = a or c or g or t
caeggecage cteteetgea getgegegtn geteaceteg etetggecee tggtgeegte 60
cacctccagg gtggcctcac cgtccctcag cgagacggtg accacgtgct cttggccgtc 120
gcagacttga tetecattag ggccaaggeg tatgetecac ggccaggace accagetget 180
tcttgagttt cttcgtggag tgatagtcta ccagtgccac agagagaggc acggcacgga 240
qqtcqqqggc ccaqanqcqc aaacaaqcac qcctgtgtct qcqqctgqqc qqattgtqaa 300
gccacgactt ctacttccca ggttgattca gtcccgacgt ccagaagggg tccgcatgta 360
gtccaggctg tagaaggcga agcttncccc ggggttagaa agaagcctct ctccgtcacc 420
gagaagcact gcatcctcgt gttnatttca ccgttttcct ggatggtggt gtcttctccg 480
ttcagccagt t
<210> 36
<211> 436
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA070752
<220>
<221> unsure
<222> (1)..(436)
\langle 223 \rangle n = a or c or g or t
<400> 36
acgtgcagtt cagtcaatga aatcctgagg attggataaa gtaaacaaac tgaaatggat 60
gcatcgtacc atctactgat gaggaagata tgaggtccta gttgtgaatc atgaaatatt 120
tagagtctgg gtacccatga gttagaagag gatttgctga ggtcatttag gtcttcattc 180
tgctgtgatg tccagttgag ctactgacgg tcctctggct gcttctggaa actgatgctg 240
gcataggcgc ttaaatcctc acttgagcgg cgggtggagc tgctctcacc gctgcccagg 300
ggttgatgan ngggtggggg tgggggaagg ctgcggttca ggggtgcact cctgagggca 360
ctgtttgaag teettgacca aatecaggte tatgtagtta agaccattet ecaaacece 420
agcagcccca cacagt
                                                                   436
<210> 37
<211> 567
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. AA082546
<220>
<221> unsure
<222> (1)..(558)
<223> n = a or c or g or t
<400> 37
agagaagacc gtggatcacc tggggacaga ggtgaaaggc ctgctggggt gctggaggaq 60
ctggcctgga acctgcccc gggacccttc agccccgctc ccgaccttct cggagatggc 120
ttctgagccc tggagctgga gcccagcagt tggaggtggt gcacctgcca ggcagcgcca 180
cagaaccagc cetgteetet egaetteett cettagette atgtgaaata aaagetatte 240
tggtctcctc tgtgtctgct gacagagtaa cccgtttaac tacagcctcc tctcactcca 300
```

```
cttccatgcc tggaggaagc ctgcaacccc ctccaggctc agacctgggg acacccccan 360
tcctgtcatt tataggggaa gatggagcag gggttgattc acacagatgg ggggccctct 420
gaattggcct gcttctcaga atgttggcca taggtnaaaa gcaaggggat cggggttcag 480
gaccancaga atgtttagtg aatctgnatg aatgagaccc caggatttat gtgtccatta 540
agtggttgtt gtgntttaaa aaaaaaa
<210> 38
<211> 328
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. AA084138
<400> 38
ggttacaaga ttctttattt tgtaaactat acataaacag taaaaaagaa aatgcattat 60
actitattac gtaaagtcaa cattaaatti tgtattgagt qtqtataaat taaatqqaaa 120
taattaatca attttgcttt caatgaattg tatactggga aaccagttta cccactgttq 180
aaattaaaga taccaatacg taacattcaa caggtttttc catttttatt atgggcacaa 240
aaccattggt atgatatagt taaaagtgat ggtgtgccaa aatgtctaca caattaatta 300
acatgctaac ttaaatacag cggttaaa
                                                                   328
<210> 39
<211> 370
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA085943
<220>
<221> unsure
<222> (1)..(370)
\langle 223 \rangle n = a or c or g or t
<400> 39
agaacccagc ggtgttctga ggggagcgtt tatttcaagc naccgatggg acaaacantc 60
ccaggettee caggtgnean tgneeggge ggeateetea ettecagegg cetecaaege 120
ggcccttccc tgccccttc cggaacttct gggcgtggct gatgcggttg tacagcacgt 180
tgateteata tttetgetgt tteagetteg ceateaggte gaaettetea gaeteeaget 240
ggtggatcca gtccgacagc tcctgggctt tctcccggag ctgttcctcc cccatgtaag 300
tcaatgttca agagggcttc ttaacgctcg gaaaaggaat gcgcaccttc atctcccggc 360
ccccgtctgg
                                                                   370
<210> 40
<211> 406
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA086264
<220>
<221> unsure
<222> (1)..(406)
<223> n = a or c or g or t
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<400> 40
ttatttcngg aagtcagaag aaaaacaang ngcacaacct gaatgacaca gagcggcagn 120
tggaaaccac aggggctgcc ganagctggc ctttcacagc agaccactgt tttccagtga 180
gaatggtggg ccattccaaa acaaagctaa agggttccaa acatccagaa tggaagctgc 240
ttcccccaac tccattacct atactacagg atggattgct ttttgtgaga ccccttcttc 300
cactgggcaa ttttnggcat tatttaccct cccccqatt tttaaaagct aaaatggcgt 360
cccagggaag aagtgccggc ttggatgcan gcttgggcca ntcact
<210> 41
<211> 250
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA091278
gtttgccttc taattgatca tttagactat tctggctaag tctgcccaca tgtaattacc 60
ggctaattca agcgaggaaa aatgtaagtc atttagacca aagccaagca gtttctttqc 120
gtgggttact caagggcttg tggttacttg tatctcctct atgtgaactt gactttgaaa 180
gacagagete tagtgtgcca geetgetaag teetgtaaga atagggaggg eggaggggt 240
ggcagtacta
<210> 42
<211> 307
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA092716
<400> 42
gcgagtctgg aactctttct tcggggcccc ggggcacacc atggaggtct cctgttgaat 60
ggcccttgtt gccctagagt gggacccagc cctcacctcc cccagagcta acctgggagg 120
tgctgaaggg gcattgggcc accgtaagca agggaaaaag ggcagatcat gcggggagat 180
gacettgate tttgattgct accetaacet tgacetttaa ceegtgatte ceecagetee 240
tggagagatg tctaatatct cttagggacc agaccctaaa ttctctctcc ccatttgatg 300
ttagtgg
                                                                307
<210> 43
<211> 309
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA093923
<400> 43
gtcataatgg accagtcatg tgatttcagt atatacaact ccaccagacc cctccaaccc 60
atataacacc ccaccctgt tcgcttcctg tatggtgata tcatatgtaa catttactcc 120
tgtttctgct gattgtttt ttaatgtttg ggtttgtttt tgacatcagc tgtaatcatt 180
cctgtgctgt gtttttgatt accctggtag gtattagact gcacttttta aaaaaggttc 240
tgcatcgtgg agcatttgac cacagtggac gcgtggctat gcaggtgatt cctcagtctt 300
ccttggtct
                                                                309
```

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<210> 44
<211> 271
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA094800
gcgactgcag aaaaagttcc agaaacaatt tggggttagg cagaaatggg atcagaaatc 60
acagaaaccc cgagactctt cagttgaagt tcgtagtgat tgggaagtga aaqaggaaat 120
ggattttcct cagttgatga agatgcgcta cttggaagta tcagagccac aggacattga 180
gtgttgttgg gccctagaat actacgacaa agcctttgac cgcatcacca cgaggagtag 240
aggccactgc ggcatcaagc gcatcttcac a
<210> 45
<211> 323
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA099820
<220>
<221> unsure
<222> (1)..(323)
<223> n = a or c or g or t
gtgacatgtt ttttgcttta ttgaaattct ctcttacaaa aggtctgang tattttaggc 60
caggectaat ttgetttggt ceetgaaatg caggeceatg gteattteea tgteetetga 120
agtaggtatg taaactagta gacttccatt tttaaggttc acacactttt taacattqtt 180
tttatttgat gtaaaacaag acttatgttg tccctaatgg aaagaccaag taagagagtt 240
atgtgcgtct tcatggaagg gataactgga ttctttgcca gaaccgggtt gggaatttag 300
tttgttcaat gtggcatctt tca
                                                                   323
<210> 46
<211> 431
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. AA101767
<400> 46
catttcataa ataatgtact ttattttatt gcatatggct attaaggagg gcatccatga 60
tcaatacaga ctaaatacaa tgcactattc tagtccagtt tattctcgtc tccagcagca 120
tcacattgac ccctatatac agcgtgtaca gtggaagaca gagcaagata agttaagtct 180
cttgtcatat cacaatagca agaaatatat ttaacatctt gatatccaga aacaatacgt 240
acccaaaaag aaaacactgt ttaataactg ttaaagttta tatagcaaaa aatattttaa 300
atttaaggta agtcaggcaa aatgtacaaa gacccaatat acattgtgaa gttttaqcaa 360
acataacatt tatacatttt ggttccattc tgtaaactaa attaaaaatg gtaaatattg 420
catatgcctt t
                                                                   431
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<210> 47

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<211> 260
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA102489
<400> 47
agtetacaag ttcagaccca catgtaacgg atttttgctt catggttgtc agaggctagt 60
gtgcattatt tctgaggatt atatccaatg acacgacgca gaaaacacaa atggacggac 120
agacggatgg acataatcat taagacaaga gactctaaaa cgtgccttag tgtccacgtg 180
attgatctaa ggcggggacc cttctaaggt ggggacccga gtgatctaaa gcagggtggc 240
ttccagcaca agggtgccga
<210> 48
<211> 365
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA121142
<220>
<221> unsure
<222> (1)..(365)
\langle 223 \rangle n = a or c or g or t
<400> 48
tttttttttt ttttcaacaa actcagcttg actttattac atggaagctt gcagggagcc 60
ggtgttccag atctgccggg gagaccagat caacagcctg cctcttcagt ttatatccgg 180
aagactcqcc caqqtcctqq ctacttqqqq ccaaqqtaqq aaacaqcctt tcctqttttq 240
ttgagggttg ccancagggt gtctgagctg tgcccaaagt cgatgcagac cttctttttg 300
ggcaaggtca atgttgaact ccantcctcc caagcttgtt tgaaggactc tggaaaacgg 360
gtttt
<210> 49
<211> 261
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA127946
<220>
<221> unsure
<222> (1)..(261)
<223> n = a or c or g or t
<400> 49
ttaaagtgaa agaaacttta ttttgagtaa tatacatatc attcattcca tttaattttc 60
atagctatgc nctatgaaaa ttaaatggaa tgagtaatat acatatcatt cattccattt 120
aattttcata gtgcatagct atgtgtagaa gtacacaggg aagaataaac attaqaaata 180
cctagccatg aaaatataca agtgaagaca tttgatatat ccatggacng gcttggaagt 240
attataaaac aggatccatt a
                                                                261
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<210> 50
<211> 444
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA130349
<220>
<221> unsure
<222> (1)..(444)
\langle 223 \rangle n = a or c or g or t
<400> 50
tacaaaaaac aattgttatt tgtgtacttt taaaacctca cagtaatatt ttcacactac 60
cttcttggct gaaagttcac actcggaatt ccagagcagt ccatggccag gcccactgqn 120
teccettget eteteettgg etttggtaac caetggeece agggaeteag eetgetttee 180
tatccatccc ctcagtagct gtcaccatgc aggttacccc ttctgtttct tctaccacta 240
actccatgtc tgactgcaag tgaaaggaac agaagcccaa acctttgggt tttaaggagt 300
ttattgctaa tctgtaaaac agaaagagac aggagataag catgacaaaa tatagggaag 360
aaatgacttt tgcctaaact tccaaactgt gtacaattga agcctccgct ttatagctct 420
tagcacacct ctcaaataag aagg
<210> 51
<211> 616
<212> DNA
<213> Homo sapiens
<220>
<223 > Genbank Accession No. AA131322
<220>
<221> unsure
<222> (1)..(616)
<223> n = a or c or g or t
<400> 51
gatttccatg cactttaatg aggtccagca ctcaggagga ttagcqccca ccaccaqctq 60
cctgggcagg ggagggccgg agcaggtngc aggcgtcagg cttaggacag ggaagggggc 120
tcaggatggg gaagggtcct caggacaggg gaaggggctc agaagagagc agggggctta 180
ggacaggaag gggcactcag gacggggcag ggaaggtgtg gggggcagtc gccacctggg 240
taggaagcag tggtgttttg gacaggaggg gctggctctc cagtgaccca ggtggacacc 300
ccaggcctga ctcacggctt tttggggaca tagtggtgga tccagtccaa gtagtaggtg 360
acacgggtgt agatgccagg ccggttgggc tgggcacagc tncgntccca gctgaccacg 420
cccgcctgta gccaggtgcc attcaccttg cacaccaggg gccctccaga gttcgccctg 480
gcatgagtcc ctccggtgtt cccggcacac agcatgtcgt tcacggatga tgccgacgtc 540
gtctcccgtg taggcgccaa agtggtattt gcgtcacaaa tgtggtttcc attatgggga 600
ccttcactgc ttcagg
                                                                   616
<210> 52
<211> 464
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA131919
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<220>
<221> unsure
<222> (1)..(464)
\langle 223 \rangle n = a or c or g or t
<400> 52
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ctcaagtgat cctcccactt tggtctccca atgtgctaga attacagccc tgagccacgg 120
ccccatgccc cgtttttacc agtgtatatt ttctactgga aaatgagact tttagggatq 180
aatgtggact tgtctgttga aacttgtaaa tttqcttaaa aaaaaaaaqa tctccaaqtc 240
ttcacaaaat tttatattcc ccaaggctgc cccatcacaa tgcctgtgaa gcttgactgg 300
cagacactga ggcctgaagc tgggggctgc agggggtcac tggctcaccc ggtccccccq 360
taatctgtaa aacatactgg gtgagggagg ctgctggagg acctgaatct ctcccttctc 420
caggcagtag tgaggcatat gctgntggcc ttgggccaat taaa
<210> 53
<211> 393
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA133756
<400> 53
tacgaagatg caacaaaatt ttaaaaaaga aaaaggggtg caatttttt cagagaggac 120
agctgatcaa atatttataa ttttctaaac catgcagttc attacttatt acaattccaa 180
acaaaactca ttattatggg gatgggagtc agggagaggc cccccccaa gcatgatatc 240
cagogotgtc acacagtgct tatgttcaaa gtgcttacaa atggtgtctt cacaqcataq 300
ggaagctgaa gccttattcc agggaaggag aggtgagtca gtagcagtgt ccaatggcag 360
actcagaaag ctcggcagtg acttgctcaa aat
                                                                393
<210> 54
<211> 398
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA135870
<400> 54
aaaatttaaa ataaaatttt attttatctt atactcaagt tcagacaata gcatgtggtg 60
tacattcaaa atttttgaca ggtacagagc acattaaaaa atgaagacat gatcaaggag 120
atgtaagaga caaatagaca acaacattct ccctgaatct ggaaaaaagc aagcaataag 180
atcacgaaag gcagctgtaa aacaggatta ttctgcatgt gttgcccaca actagggcaa 240
ggttatctct catcacaagt acaaagccat tgatgttagt gtgtaacaga gagaaaacag 300
aggatttgta cagctgagga aataaatggc agatgttaca caggaagcaa tataacatgg 360
tcattaagta actgtattca accctcaaat ttaatttt
                                                                398
<210> 55
<211> 390
<212> DNA
<213> Homo sapiens
<220>
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<223> Genbank Accession No. AA135929
<220>
<221> unsure
<222> (1)..(390)
\langle 223 \rangle n = a or c or g or t
<400> 55
aaagatatca attatatatg tatataaaaa aaaaacctca ctttccccac aaaaagcaca 60
atactgttat cacaaaaaa atcatcatcc tcataattaa tcatcctagc cacgcaggtg 120
tntttgctgc caaaagatgg gacgacaaat aacgttgacc aggcagaacc cctagacacc 180
ctcggcccac ccacagcctc tccggctgcc gaagacgagg gacgagggca aggcagagtt 240
ctctgaggtc cccaggcctt caccccatct gtcagtctgt gtcttctagg acagaaggta 300
gttgtttttt tttcttttaa aacgtctgtt caaaataaaa aacaaaagca cacgcgcaag 360
agaagcgggg aggaacggag gctgcctgcg
<210> 56
<211> 511
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA142858
<400> 56
ttttttttt tttttttca aggggaaact ggggcagttt tattgacgat ggcaatgtac 60
aagactccac acctaggtat gtgcacgagg taaggcctga gctcaggcct tatgatcctc 120
ctcaggaccc ttgggggcaa acttctcctg cagtttcttc cacatgcctt tatctatttc 180
cttaagctct tccaaggtgt ctgtggacag gatcagcttg tactcttcca acgacaggcc 240
actgaagctg gtgtctctgg ggcgagggta cttgtgtttg tagtagtttg aatggagtcg 300
cgctaagtct cgtacatctg atcacaggcc tcaggtctgc aacctgggta ttctctccct 360
ccegaaaggc ctgtgctacc cgctgtcgca ggtaagcgcc caagtcccgg ccccgtttgg 420
tetegteeae tggeeattee teacagaget taagaaaaeg ceggtaeegt gggeegeeat 480
ttgggcccg cgtgttcccg ccctcgtgc c
                                                                    511
<210> 57
<211> 341
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA147224
<220>
<221> unsure
<222> (1)..(341)
\langle 223 \rangle n = a or c or g or t
<400> 57
aatacatttt cacagtgtgc tgaatgtctt tatttacaag atatcattct atagtgaata 60
tgaacaaaac gaatgtgcat ggttgaaata actgcttgat taaaaatgtg ctgtgaagat 120
gaatcactaa tetttetaat geactetgat aacacaataa acatggaaaa atactaatee 180
cctaatagat cnaaatatag natatagncc ccnaaatatt tcngggggat ggattttcct 240
tengaggttt encaaaaagg naaaanggaa atggntteee ceagecaatg gtttagecaa 300
atattggggg aaatgcccat tccaatggga aaaacccgga t
```

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<210> 58
<211> 561
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA149579
<220>
<221> unsure
<222> (1)..(561)
<223> n = a or c or g or t
<400> 58
atagtaaata tattacattt attctaaaac ttcaaaatta ttctgttttt gtagtactga 60
aaaaaagaca gtgccatttg aaacaacaga tgcatctttt atacattttc acaagtttgt 120
ttttcatatt tttaaaggcc ccatttatct gtaacagtgg tatttttatt tagagtatcg 180
gctacttaat atatacatgc aacaatatat gctttaatag tcatttaact tttaggaata 240
tttcatcaca ttaagtggtt aagcatagtg ttaaaagagt ggaatttaag gaataagaaa 300
atattgaaaa tacgctgtta ttttcatttg ttcactataa tagaatgttt ttgcccataa 360
aagttatcat tgcccaactg aattcctacc aagaactaac aagtgattct cagtgggag 420
aantttnttt nntnngaata tagagggctc gttagaaagt gcagatntag gcgggcgcgt 480
antcacaccg taatccagca cttggaggcc aggcgggcgg tcacgangta ggagatcgag 540
accateegge tacaeggtga a
                                                                    561
<210> 59
<211> 420
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA150920
<220>
<221> unsure
<222> (1)..(420)
\langle 223 \rangle n = a or c or q or t
agcgttgtaa ggtttatttg ggtagggaag gggacaagtg aggtaactga tccttgcttt 60
gtagacagtg caagacaatt atttgtggtg aagggactgt atgccaacaa acgttactca 120
tgctttagtt aaaactttaa gtcacctaaa acagaaacaa ttctnaagaa cactggtgga 180
aaatagaagt gtaaatgttt cagacaaaac caaggcattg tcagcacgat gtacattata 240
cggcagatan nacagccaca tcctaggcca cagagcagat cccaagagcc ccaggcatgc 300
aggagagttt taaaggaaca gacggaaatt ttaactgtga aaaccacgaa atttcatgac 360
ttttggtcag ctacnacccc aactaatata tgaccattaa gagtaaaatt ctgaccttta 420
<210> 60
<211> 426
<212> DNA
<213> Homo sapiens
<2205
<223> Genbank Accession No. AA151210
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<220>
<221> unsure
<222> (1)..(426)
\langle 223 \rangle n = a or c or g or t
<400> 60
tttttttttt tttctggatg aatacatgtt ctggtcttgt tacaggttct ggtaaatcag 60
atggaqaaat gttgttqcaq aaatgtcaqc aaactttaca qcaqtaqttc acacatgcag 120
ctactataca ttcattcatt gctattttcc taagaaatgg agcaacctag gagcttatgc 180
tacagtagat tecaatgaac cataatgact actteaagaa caaagaagca catneaaagg 240
tgtgatatct tcctgttggt ttgagttttc aaacctgaaa ttctttaaaa tacatttctg 300
ggattttatt taaatattga tgcnacacac ctaaaaagca gtgacttctt gggtaaaatg 360
taatactgaa atggaaaatt gtcttttcaa aaaaataaga agtgtggttt ggaaattccc 420
cgtgcc
                                                                    426
<210> 61
<211> 400
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA151428
<220>
<221> unsure
<222> (1)..(400)
\langle 223 \rangle n = a or c or g or t
<400> 61
cagagagaaa gtgctttatc agccgggctc agcccgcaca cggactcgcc aggagtaggt 60
ggtcagcacg cgctgctggc ggcnaccacg caggtgtagg tgccctcatt gacggcgttg 120
gcgatgatgc tcaggtgcgc ctcgcccagg gccaggtagc cggggtagga gaactccagg 180
ggctcctggt ccttgtacca gtacactttc cctttcttgt ggaggatctt ctggccgcag 240
cggaaggtca cgttcctgcc ctcggnacca agcctggttt tggtcctggg gggcggtggn 300
ggtggttggc caccgtgggg aaaggggaat ttcgtagcaa gaaantccgc aagctngctt 360
gggggcaaaa agcttccttt ccantgaagn cccgccggga
                                                                    400
<210> 62
<211> 502
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA151544
<220>
<221> unsure
<222> (1)..(502)
<223> n = a or c or g or t
<400> 62
caggacgagc tgtgggggct gcaccggctc tacggatgcc tcgacaggct qttcqtqtc 60
gcgtcctggg cnggagggc ttctgcgacg ctcgccggcg qtcnatgaaq aqqctctqcc 120
cagcagetge gaettetget acgaatteee etteeceaeg gtggccaeca acceaecgne 180
ccccaaggac caaaaccagg ctggtgccga ggnaggaacg tgaccttccg ctqcgqccaq 240
aagatcctcc acaagaaagg gaaagtgtac tggtacaagg accaaggaaq cccctqqaag 300
```

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ttctcctacc ccggctacct ggcccttggn cgaaggcgca ccttgaagca tcatcgccaa 360
cgccgtcaat gagggcacct acacctgcgt ggttgcgccg ccagcagcng ttgctgacca 420
cctactcctt ggcgagttcc gtgtgcgggg ctgagcggct tgaataaagc aatttctctc 480
tgaaaaaaa aaaaaaaaa ag
                                                                   502
<210> 63
<211> 285
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA152200
<220>
<221> unsure
<222> (1)..(285)
\langle 223 \rangle n = a or c or g or t
<400> 63
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tagetteeeg geeteaegag tgttgaatga catgaegaat teteetteat agaaggtaca 120
ggtgaaccag aactggaggg gcatttggga tccttccttc ttcagaaagt gcgatcgcat 180
caagatgcat gtggttttca gtagaactgg cccatgtttc ttgggagcga ggtgtccaaa 240
ccactgttca tccatatttc cnggatgatt tgctcccngg gctca
<210> 64
<211> 457
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA156565
<400> 64
atagtaaata tttaattgtt tccatcagca attccagcac aagttttcct ggatggtagq 60
cagaatcaag ctacccaagg gttcatgatg aggtatgggg gtcactgagg agacccccaq 120
agtcactgac ccctcccgcc acctccacac accaggtggc cctgcagaat gagggttggg 180
ctgatagaat gtcaattagg ggagacagga tacagggtga gggaacaggg tctagcttgt 240
atatttgcct gcaggaagga gggagggcag gagagactct gcatagaagg actggaacta 300
cacatttaag ttttcaaccc caatatgcag ggggaaacag ccaagccact ctccatctgt 360
ctagtattag gaacctctct tcaagtggtc ttttgtcatc tctgttcttc ttcccaattc 420
tgtattccag attccaaatt ctacaattga aacccaa
                                                                   457
<210> 65
<211> 428
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA156897
<400> 65
cagacatgga aatataattt taaaaaattt ctctccaacc tccttcaaat tcagtcacca 60
ctgttatatt accttctcca ggaaccctcc agtggggaag gctgcgatat tagatttcct 120
tgtatgcaaa gtttttgttg aaagctgtgc tcagaggagg tgagaggaga ggaaggagaa 180
aactgcatca taactttaca gaattgaatc tagagtcttc cccgaaaagc ccagaaactt 240
```

```
ctctgcagta tctggcttgt ccatctggtc taaggtggct gcttcttccc cagccatgag 300
tcagtttgtg cccatgaata atacacgacc tgttatttcc atgactgctt tactgtattt 360
ttaaggtcaa tatactgtac atttgataat aaaataatat tctcccaaaa aaaaaaaaa 420
aaaaaag
                                                                   428
<210> 66
<211> 602
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA158262
<220>
<221> unsure
<222> (1)..(602)
<223> n = a or c or g or t
<400> 66
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gatcactcat tgtatccttc tccacctttc ttttcttctc ttggggtgga gcagcacttc 120
tgactgtccc tgctgactga gcttttaaaa cttctgtaga ttcctctttt tcagttttct 180
ttccagcagc tgtaggcgac ccacaggtga agtcagatga caaggcgtct atagcatcat 240
ctggccctat gggtttagcc aatagttccc tatattttgg aggaattgtg acttctcttt 300
tacccaattc ctctatgtag gtggaactca ttggatctga aacttctggt ccaqtatacg 360
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cagcatccat gcctgacttt cccgatggtt tatccggttt agattcaact ggcacagctg 480
gggttaatga tttcttttct tttttcttgt canceggctt gcagatattg cagtgatacc 540
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tt
                                                                   602
<210> 67
<211> 392
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA159025
<400> 67
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atagatteta aagaatatgt ceetatgeae ageeeteeet eeceaaaaat aacqetqqqq 120
gtaggcattg cctttccccc ttgggctcct cgggtqtatt taaaaaaatg ttttqqcaqc 180
tcagtgttta tcatctgggc atgggacacc atgtccatgt ccccatattc ctagggtaca 240
gcagcagtag atggctgcaa caaccttcct cctaccccag cccagaaaat atttctgccc 300
caccccagga tccgggacca aaataaagag caagcaggcc cccttcactg aggtgctqqq 360
tagggctcag tgccacatta ctgtgctttg ag
                                                                   392
<210> 68
<211> 476
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. AA165312
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<220>
<221> unsure
<222> (1)..(476)
<223> n = a or c or g or t
<400> 68
tcgtnncntc ggttctgaga aataggcact ggcaatttac acatgccttg ctgtgtaatc 60
tcactatatt tqctcaqqca aaqtqqqaqa aqcaqcctta qqttttcatt ctaqaqatqc 120
cggctttccc acctgatcgg cttagagttc acgattgact gttttgggct tcatttcacc 180
ctctacataa caagcgggtg gactagatgc cttagcaaqq gtccgtgttg tgtqqtqtct 240
ccagccacgc actcagctca atcttagcac agttaaaaaa tqcctttcta gcaaqttatc 300
tgcccagtgc ctgaaaaagt atcatttctt gtgttcaata aaaaagcctc ctaatttaat 360
caaggaccta tggagataac tgtcttttag ttgtggcatt gcaaggatac aaatgcagag 420
atattttaaa agtgatcctt ctgtaagagt gaacccacga tatgatctgg nagcaa
<210> 69
<211> 479
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA165313
<220>
<221> unsure
<222> (1)..(479)
<223> n = a or c or g or t
<400> 69
cacaagcccc cacgtccata gccaagtttt ccccggtttc ccagcagcca gtgacttctg 60
tagcattagg attettatag tagttattgt etacatttet cagcagattg aatatgtact 120
gcctcttact actggactgt ttattcttaa atgtgtacag tatggattta tgtcgtctat 180
atattatgca tttatttgtc ttcttcgttg tgatggtaag ctcctggagg gcaagtcttg 240
catccactgc tttgctggca acccgactgg taagcttctg gaaggcaagg cttgcatcca 300
gtgctttgct ggcaacccga ttgctaagta ccgtgtttta agcttagttc agtctcaagt 360
gtttgcagcc acatctgaag accaataaag caactgctgg gtttatcccn tgggagctga 420
cagaatttcc tctcccaaat accatanaca ggaaaatcat aagcctgaat tacccggtg 479
<210> 70
<211> 298
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA171939
<220>
<221> unsure
<222> (1)..(298)
<223> n = a or c or g or t
<400> 70
ttttttgagg cacctgtggg actttattag gtaaacagac cccagctcca gccacaggtt 60
ggaccggcca gctgacagtg cggcctcaga caccccgcc aggttccctc ctcctcctc 120
tctcagggtc accagtgtgt gaaagatcgg qqcatqccqq ccacaqqggg aaqcaqqqtt 180
caggetgeec cacetgggte tggceetgge aggegeece teacetgget etgetgtggg 240
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298

anceqagaac aaagacatna cetgeetgge teetgetgee eeggggggte agenagea

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<210> 71
<211> 596
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA173223
<220>
<221> unsure
<222> (1)..(596)
<223> n = a or c or g or t
<400> 71
ttttttttt ttcagccaaa ttcatattta ttccagtctc taacactctg ttgttatgtc 60
tgctgtaaga tgatcaggag ttagtatgaa gtattcttct ctacgcacca aagaaaacaa 120
acaaagcaaa cttcaagtca gtgaattagt taccacagtt aaaatgcatt tgattttgtc 180
cttttccttt ttcacaagaa cgacagctga atactctttc atgtgatgcc tgatattttt 240
ctttttcttt ttctctcttt tttgagacag ggtctttaag atggggtctc gctctgttgc 300
ccaggttgga gtgcagtggt gcaatcttgg ctcattgcaa cctcagcctc ctgttttcaa 360
gtgattcttc tgactcagcc tcccaggtag ctgggattac aggcatgtgc accgtgcccg 420
gctaattttt gtattttag tagagatggg ggnttcacca tgttggccag gatggtctcg 480
aactcctgac ctgaagtgat ccacccgcct cggcctccca aaagtgctgg ggattaccgg 540
tgtgagccac tgtgccagct ctgatggtga aaatttcngg tacaggccta gcccan
<210> 72
<211> 408
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA180314
<400> 72
ttagcaaaaa cagctttttt attgtggtag tttgtggtat gtgctcctgg atcatgcaga 60
aaaaaggctg ggcctcagtt agctccggga gccattctta ggaccctccg gctgcacaca 120
gagaggggct gggtagctgg ctgggctggg gcacgcattc actgggctgg cacaggctga 180
ggggtctctc gcccactatc attaggcccc tccagcccgt tatgctcagc ccccggctca 240
ggatgctcca gggcgtgccg ggtatcagcc tgccagagct gcaccaggtc cgtcqqqqtc 300
tttcctgcca ggttcttggt catcatgtca gccccatgca ggagcagcag tttgatgatt 360
ttgtagcggt tgagcctcac agcgtcatgc agggcagtat ccctcgtg
                                                                   408
<210> 73
<211> 479
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA182030
<220>
<221> unsure
<222> (1)..(479)
\langle 223 \rangle n = a or c or g or t
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acaaaaacac aatgtataca ttaataaatt aagtgggcct gagtattcag tatccatcta 120
ctagaatcct aaagctcttc cccagatttc acaaaggcca atgtagatta tttctatttt 180
atcaaagttc atttgcacag ttggtgtaat tgagatacta acatttcttt tttctagtgt 240
tttaaaqata qttcacaqta tttqagttaa ttaattaatc aactgattta aatctttggt 300
aaatacaaqt atttacatqt aaaaatqttt aqctcaaatt tcagtaaaaa actggaaatg 360
accaataacc tactqccaac tqttttqqta taatccaqaa atqcatqaqc cqqactccca 420
ccattaagaa atggcactgt cnaggacctc ngatgataaa actggaatcc ncaaaaaat 479
<210> 74
<211> 313
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA182882
<220>
<221> unsure
<222> (1)..(313)
<223> n = a or c or g or t
<400> 74
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tecttgeete tgeeetttte accteecet eceteceage ttettetgee tagagegtte 120
cagattecce teacatttte etggateagg gecaeteete ceaggeacet ettgeeetea 180
ccagtacett ttgtcccttc tcctggggct gagggtcctc agctgtgctg gnccccaact 240
ctccaccctt agtgcccact gtctctgcca ccctcccttt ggaactcagg gggctcaggc 300
atcctggcct ctg
                                                                   313
<210> 75
<211> 258
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA188981
<220>
<221> unsure
<222> (1)..(258)
<223> n = a or c or g or t
<400> 75
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tcttcattct tcagaagact taattagagt agctttcttc tcatacttat ctctaatctc 120
tttaatattt tccgagagat cttctgacat gcattcntca tattctctat caactttagc 180
aatctgctcc tcaagatgtt tctctacaga cccaacatgt gtagcaacca tctctaacag 240
acgttgcaag ttaatttc
                                                                   258
<210> 76
<211> 506
<212> DNA
<213> Homo sapiens
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<220>
<223> Genbank Accession No. AA189083
<400> 76
ttttttttat tccaaatgtc tttattgaaa cagaatgata gagcaagaaa taatgaggtc 60
tgggtggatg tetttgggeg caggatggag cecagaceca gtggttacag tgtggagete 120
tetecetgte eeetgaetet ggeeaaggaa gtgaatgeaa ageageaggg aggaggeagg 180
gtggggacgg ccctctgagc tctccgcgat ggctggcgtg aggtgcctct gagacttctg 240
ggcagccctg cettccctac tcagtettcc cgatettett gccacetttc tgtgtgggcc 300
agcctcccgc cagtaactca gaggccgctc agagggcagg gttgggggtg gcaagcagcg 360
ggacgtggtc acagcgggta gggggtggct gccgcagcag qqaaqqccqq cqacacaqct 420
ccccgtcccg gagcacctcg ggcaggagct tgcgcttggt ctccggaagc agcataatgc 480
tgaaqaatqc agaaqaqqc qcaaqc
<210> 77
<211> 513
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA193197
<220>
<221> unsure
<222> (1)..(513)
\langle 223 \rangle n = a or c or g or t
<400> 77
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ccatttatta cacttctgaa gtaggatttc tgaaqtcatc ttatqqcatq taattcttaq 120
tataatgcac aggattcctg tcattttgaa gcacgaggag aggtttttga tatcttaaac 180
attittttag tgtagatgca catattctcc acttccaatt gtaatagaaa atcagtttaa 240
ggatacccta atgatgcaaa tgaaatgatt agcaaacaac tcaaatttag gagccttctt 300
ttattagaaa atggttttaa actctgatca ttacattgaa gagtcaatga ctgaggtttt 420
cttacctact ggctcatctc ttagacaata acttcttgaa taatttcnac atgagtgtct 480
gtacaagctt ttaaaaaacc gaataaatta aag
                                                                 513
<210> 78
<211> 499
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. AA195678
<220>
<221> unsure
<222> (1)..(499)
\langle 223 \rangle n = a or c or q or t
<400> 78
gaaaatttgc ctcctggtaa ccctgtaatg gatggggccc agaaatgaaa tatttgagaa 60
aaacaagtga aaaggtcaag atacaaatgt gtattaaaaa aaaaaagcct attaataggg 120
tttctgcgcg gtgcagggtt gtaaacctgc ntttatcttt taggattatt cctaaatgca 180
```

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tcttctttat aaacttgact tgctatctca gcaagataaa ttatattaaa aaaataagaa 240
tcctqcaqtq tttaaggaac tctttttttg taaatcacgg acacctcaat taqcaaqaac 300
tgaggggagg gctttttcca ttgtttaatg ttttgtgatt tttagctaaa gagagggaac 360
ctcatctaag taacatttgc acatgataca gcaaaaggag ttcattgcaa tactgtcttt 420
qgatattgtt tcagtactgg gtgtttaaag gacaaatagc tqctaqaatt caqqqqtaaa 480
tgtaagtgtt cagaaaacg
<210> 79
<211> 463
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA197112
<220>
<221> unsure
<222> (1)..(463)
\langle 223 \rangle n = a or c or g or t
<400> 79
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aaaatcctga agaaggtgca aggtgagacc cagtgcgagg ggcgtgctca gatatgcagt 120
gtgtgtgtgt gtgtgtgt gtgtgtatcc gtgtgtacat gtgtgcacgt gtgtcgtatg 180
acgtgtggcc cacagagggt ggggagaaag cttggctttt tacttccatc caggagggaa 300
ggagggcggc tggtcctcca gccttggagg gtctgcagct gggcgggacc tctactcagc 360
caggetgttg cgcatcgact cetteteetg gagggeggee atggeaagae geaggtgete 420
cttcagctgc tcgatctccc gctcagaccg tgtctngatg tga
<210> 80
<211> 404
<212> DNA
<213> Homo sapiens
<220×
<223> Genbank Accession No. AA205376
<400> 80
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atacaaataa gttcattaaa aacacaggct gattattcat atctattaca ttcaqaatta 120
tgcgaaacaa ttagttatat tgcaaagctg taattctttt tctaacaaag catgatttta 180
taaaaacttta atgttgccac tgattcaatt ttaatacaaa atacttatat acacaataca 240
atataaaagt aaactgtgta gtgccttcca caaagggata tattaaggcg ctttacaaat 300
ataccaatat tttgacccaa attacttttt gctttagatt aaaatgaaca ggctaaatgt 360
tccactttaa ataccaaagg gatggtttat taaaaatttt ttat
<210> 81
<211> 523
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA205724
<220>
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<221> unsure
<222> (1)..(523)
<223> n = a or c or g or t
<400> 81
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tggaggtagc tgcaaqgaga atqtctcttt ctcatgacaa ccaaaqcqac caaaccatac 120
cctaaagcaq aqacgcaatq gaataagtca acqqqcattq taqaacqaca ctcaqaaqca 180
ggaaaaacca taaaagatac aqqatgattq tctcttcaqt attqcatttq qccatqtatq 240
tgtttttaca taaaatatat gttttctttt taagctagct aaagaaaata ctcttgatcg 300
gggttagttc ttaaagcaaa aaacagaaqa aaagtatqta tatataatan aattaaagaa 360
cgatagcatg ttatacctgg aaaggaccgt gggcactaat ctgcactttg ttccaggtaa 420
tccatggctc tgagagtgag cacactgtca aagtcactgg ggtgagatga gccgggactt 480
ggaaaaccct ctcttaactt tcagtctcaa ctcctcccac tcc
<210> 82
<211> 587
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA211443
<220>
<221> unsure
<222> (1)..(587)
\langle 223 \rangle n = a or c or g or t
<400> 82
catttagtca aatatttatt tgaactcata caaagtttag tgacataatt taaaaggtga 60
agaactaaaa cgcattccaa atattgacca aaatactgta ggaagtagct tgggaaactt 120
ttcatcaaaa tcgttaggca cattgccata tcattctcca taaaatcata tccctcctca 180
aaaccacacc ctccaggtgt tgaatttatg ggctaatttg ttctgtgagg tgccaaaaat 240
gaagataaag taagaaatac agccaactag aaggaagaga tataaatgta caaacaggcc 300
atttctgcta gagtctcagg cattcaggag gttcacaatc atcatacaaa tatataaaat 360
tttagtgagc tattgaatcc atcttctgcc tctttatttc ttcacatcaa tccttttttc 420
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ccgatagatt gactgcaggn ccgggaggtt cctcgctccn ggaattggct tcttctcctc 540
atccgaggtg ggaggacacc ctcctccact tcgggggaca ttctttt
                                                                   587
<210> 83
<211> 382
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA214688
<400> 83
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ggggcccatc tatcctggac acggcatgat gttcgtccgc aacgattgca aggtgttcag 120
attttgcaaa tctaaatgtc ataaaaactt taaagagaag cgcaatcctc gcaaagttag 180
gtggaccaca gcattccgga aagcagctgg taaagagctt acagtggata attcatttga 240
atttgaaaaa cgtagaaatg aacctatcaa ataccagcga gagctatgga ataaaactat 300
tgatgcgatg aagaqagttg aagaaatcaa acagaaqcqc caaqctaatt tataatqacc 360
agtttaggaa aataagagct ca
                                                                   382
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<210> 84
<211> 398
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA216589
<400> 84
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ccatctgcat cgtcttccta caaacagttt ttcttctact attcggttat ttctcctttt 120
tttgtttcct atttcagaat caaatttatt ttacttgcaa agtcagtgga atatggtttg 180
gaaccagtag ggcctctaac ttaagcccag aacctgtcaa agagaagtgc agtatcattg 240
ctaagacttg aacagtttat ctctcagaat cttcagttcc tttgaatttc tcagctctta 300
gtgtaatctg ttttatgtgt ttgttgtaga cttccattta tgggatagat ttccaaaata 360
attttgggta atccaactgg gtattttagc attcccgg
<210> 85
<211> 378
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA219100
<220>
<221> unsure
<222> (1)..(378)
\langle 223 \rangle n = a or c or g or t
<400> 85
ttttttttt atgcttgaac taatttattg atgagattct catttctgta gtataaaaqq 60
aaaatatttt gcagttatct cgtatttgaa agactttgcc atagagaact ttatcagaaa 120
tggatgaact tttcattatt tcttataagc atattggttt tggcctgctt gagtttaaaa 180
ctttttttgg tagacntaga atgttaatat ttagataaag aaaatatttt acngaagaca 240
ttaccagaaa gtaaaataac ttgaacattt cngtattagc ncnttatcag agaataacat 300
ttattttatt tggaaagttt tccnaaatat gagacnatcn gcnatttctc agacnaagtg 360
aaaaatttaa taaaatag
<210> 86
<211> 444
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA219304
<220>
<221> unsure
<222> (1)..(444)
\langle 223 \rangle n = a or c or g or t
<400> 86
gcttgggcaa aagtcttcag aacaaaggct gtgagcaggt gttgccctgg ttcctgccat 60
atcgctcccc aaaggtgctg taggagccat catagtgttt gtagttcaac tgtctctgqt 120
```

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aaccagtgtt gagatagcca atggcttgga cttgacctct ggagtaagct gctgtgtttc 180
atttagataa tecagtacat agatgttagg agcaaagagg accatattet getetecaca 240
gccatagggc atctggagaa gattttgtgt gttttgcatg gcagagctac atatgtctcc 300
caaaactgag acagaagctc gggcagattc ttctaccaca tttggtggca gtttcaggga 360
taattettea gaaacetean cacetgntgg acnaagtagg gagttgaatg ttgttteett 420
ctctagtcct tcaggttcaa ccaa
<210> 87
<211> 341
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA219552
<220>
<221> unsure
<222> (1)..(341)
<223> n = a or c or g or t
<400> 87
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ganggcatac aactgtcaca ggcagggcag taagtacaaa gtctagctqt aaaaaccqtt 120
tgaaaatata aactcgtttt tggaatacat gtgtcaaagg ctgcccatgt taataccttt 180
ggtataaaac ggtaacgatt cccttgacaa acccatccat cacctgacgc acattcacat 240
ctcctggtaa ctactctacc tagtctagtc tcaaccaccc ctgtcagtca cgactcactc 300
ctgttccttt gcaggtgcag aggagcctgg gaggtaggtc a
                                                                   341
<210> 88
<211> 323
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA227926
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acaaaaatgg ttataaaatg gttgaagcaa ctagaagcgt gacaggtata atacatataa 120
atacaaccaa aattcaattc aatgcaaagt tgaatgacat catattgcac caaaatttat 180
tccatacaaa agcacatgca tcaagagttt ccataagatg aaaacaaaca cacttacttc 240
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cagtaccaga actctcccca gag
                                                                  323
<210> 89
<211> 469
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. AA227936
<400> 89
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ggggagggg gcagctgtgg ggctcggcac accccgggcc ccaccccggc ctggcgctgt 120
ctgagaagag gggatctgag ggagatccag ggatcaggca ggatagggat ggggcaggac 180
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cagatggagc aggtcaggag gtggaacaat ggcagagtga gggtggaggg cgcagtgtct 360
ggagaggcgg aaatgagaag gctggggaga aagaagaggg tggcagctct ggtgcagggc 420
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<211> 462
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA232266
<400> 90
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ggaagaaccc ctccccagat tggcccagtt tcaccaqcaa ctggtctcag ctcagcctta 180
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ccaaaaaaac aaacatcatt cttagcaaca tcaattactc ttccacacaa aacagaaacc 180
ttgtaaaatt tattttcgta tttttaaggc gtaatacttc cgtataaagt atatgcaaga 240
gataaaactt cacagtattc caaaatgtca caataataat aataatata tagtataatg 300
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cagaaattaa cttgtaaatg attctctttg gaatctgaat gagatcaaga ggccagcttt 240
agcttgtgga aaagtccatc taggtatggt tgcattctcg tcttcttttc tgcagtagat 300
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gaaattcca
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<211> 375
<212> DNA
<213> Homo sapiens
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atcacccatc tgaaattcat ttacaaggtt tttacattaa taaaacagta gtgtggtaca 240
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<213> Homo sapiens
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ggtaccataa taaatcacac actcacacat ccatattgct taggttgaag agaacggaat 180
gaacagagga aatttettee atgaattgee eteetttegg taecegeeat gttttagtta 240
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CC
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<210> 127

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<211> 428
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acaagtccat gaaagtagag aggaggcgcc agttaaggga cagcaacttc aaggagacgg 180
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acacatactt tgtccggtct gaacaggtcc agggctccac cggaaactcc aatattgagc 300
ctccggttgg gtttggccta aaatttttgc ggaagaacct gggtgggcca tttcaaacca 360
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tggacggc
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<212> DNA
<213> Homo sapiens
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cogtococac coccoctoc accoctoggo coatcagtqt qtqttqqqqq qatqcttqca 180
gctgggggtg aggagacaac aaacctcggg aactggagcc agagctgcgg cctgactgac 240
gccttttgat gctcacggga aatttctgcc caggatctca gccccaggct ggttgtttct 300
acaaatctct ctcaaatgta ttattttggt gacaaaaatg aaggagcttt gtaaattttt 360
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cagca
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<211> 405
<212> DNA
<213> Homo sapiens
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<400> 129
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tattttttgt ccatcagtag ccaaatggaa cttgatataa acacttccag tatqccaact 120
ttggtttaat gcacaacttt gaaaataact cattaaaaca cacatcaaga tgctactaac 180
aaattcatta atatccaaga ttcattactg tatgtcaaag gtcatccagg attaacattt 240
tcattacaat gaactgtgaa attccaatga aaaatgtttq cctqaattaa attatttaat 300
ctctcaaatt ggaagtctag cactcttgaa aatcaaattc acacacaca agacacacac 360
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<212> DNA
<213> Homo sapiens
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<221> unsure
<222> (1)..(478)
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acagcgacag tgatgactcc aaaaaaaatg tttagaatta gaagtgcatg ttaatctgag 120
taacttaagt acagaaaaga gttagtacac cacaagcatt ttctacactt ttattttgtg 180
gtgattgtga gacaaacaca gtccaaacaa tagacttctt gtcctccccc tcccaacaac 240
tatctgactc catageteat geaceceaat tacageaggt gtegggetgg cataaagget 300
tcttaccagg attccagttt atccttctca atccttttct catctctaac aaaaatqcca 360
cacatacatg tagttgtgag aggcaaagtc ttctttacac tcaccaccaq ggnqqcqtat 420
gggagcacaa aagcctcaca aaactgctcc aggatcctqc ctcttccaqq qccqqaat
<210> 131
<211> 216
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA291676
<400> 131
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gactacaaaa ggacctaagc cttttaaact agactgtctc aactgtgcat taattatgta 180
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<210> 132
<211> 431
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA292328
<400> 132
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cctcctgcct tggcctccca aagtgctggg attacaagca taagccactg cacccqqccq 120
agaggggttt ggaatgaagg tagaggcagg gggatgaagg cgccaqaqct qaaqaccaqc 180
ccccagaagc cacacccctg cccttctagc agctacgggt cctctggctc cgggccttgt 240
aaacctcgat gagcaggtcc ttgacgtact ggatctcgcg ctccacggac tctgcccqtt 300
cetteagete gegatteegt geeteeagee cetggaacte gacceteeag ggeeteacee 360
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tgcttgcggt c
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<210> 133
<211> 318
<212> DNA
<213> Homo sapiens
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<223> Genbank Accession No. AA293187
<400> 133
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gggagtggct gttccctggg tccaccagct ctgggagggg acatggaaat ggaagatgtg 180
ggtggcattc cggacaggga ctggtgcctg agaatgctgg ggtcagagtc ctgggaggga 240
gcgagatggg ggaacatctg tgctcagaag agggggtgta tgggtaggtg catgtgcttc 300
tgtgcaaatc ctggtccc
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<210> 134
<211> 424
<212> DNA
<213> Homo sapiens
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<223> Genbank Accession No. AA293489
<400> 134
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ccccagcctg ggcccaggca accaagggct caatgctggg aaggagagca ggggaggtgg 120
gcttagtgtt aaggcgtgaa gggcgaggcc agacagctgg aggcctggtc ctccactctc 180
catttccatc accettegga ggctgaagga agggeggegg caccacaggg cccttcccct 240
ctgctgcatc atctcctgct caggctttct ctctaggcgc attggaggaa tcctctttcc 300
ctgtcggaaa ctcaacactg tacagaactc caaccataac ccttctaqct tcctctccca 360
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aagg
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<210> 135
<211> 340
<212> DNA
<213> Homo sapiens
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<223> Genbank Accession No. AA298981
<220>
<221> unsure
<222> (1)..(340)
<223> n = a or c or g or t
<400> 135
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agactgtgaa cagcttgctg tcacttcttc acctcttcca ctccttctct cactgtgtta 120
ctgctttgca aagacccggg agctggcggg gaaccctggg agtagctagt ttgctttttn 180
cgtacacaga gaaggctatg taaacaaacc acagcaggat cgaagggttt ttagagaatg 240
tgtttcaaaa ccatgcctgg tattttcaac cataaaagaa gtttcagttg tccttaaatt 300
tgtataacgg tttaattctg tcttgttcat ttgagtattt
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<210> 136
<211> 535
<212> DNA
<213> Homo sapiens
<220>
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<223> Genbank Accession No. AA308998
<400> 136
aggetetaet teaggtgetg etataatgee teatetaate aggaetaaat tgtgtaggaa 60
actgcagtgg gaagaatatg ctttctgctc aggctaagag ggtcactgat ctgtccttag 120
aaattcagag taacatgagc aaaacctcag ctaaaaccca tttaagtggc atggattgtg 180
catgatettt gataagaatt eeteatgtae ttgtgeetag ttttteaagg tattggetgt 240
tctatagatg cagtgattgt cccagctagc tctgttacca qccttttqqt qtqtctttat 300
gttcatttgg agagtcaggg cgaaagacag gtgatgtagc acttctgttt ttaataatta 360
ttgcttaaaa tacctattaa tagttttggg tcatttaaag ggacttgagg aagctaccca 420
ggattacaga agagtgtcca cctaacaaga tggtctggca gtttcctagt tttqtatctg 480
gttcaataga aatatgtgaa agtggtaatg tcatcatttg atgcagaqtc cqqqq
<210> 137
<211> 324
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA312946
<220>
<221> unsure
<222> (1)..(324)
<223> n = a or c or g or t
<400> 137
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atgcaccaca acccaattac aaagaacagg tgttaacaca caatgtttaa acaatgctac 120
actcattttt ggcaaagtgc tgtattgttc agtctgtgta caaaactgac catctatgan 180
ccaatcagta taaaaaattt ctataaaanc aaaatttagn cagtggctca agaaaacaag 240
ctgccattta tgcatagnnt gatgtacagn aacctaacca aatgtccctt ttgaattttc 300
aagttactga aaaaaaatgt gtcg
<210> 138
<211> 428
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA316686
<400> 138
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cacggaaaca tgacagcggt gcggcggact tggagcgggt caccgactat gcagaggaga 120
aggagatcca gagttccaat ctggagacgg ccatgtctgt gattggagac agaaggtccc 180
gggagcagaa agccaaacag gagcgggaga aagaactggc aaaagtcact atcaagaagg 240
aagatctgga gctaataatg actgagatgg agatatctcg agcagcagca gaacgcagtt 300
tgcgggaaca catgggcaac gtggtagagg cgcttattgc cctaaccaac tgatgcgtgc 360
tttctcaaat atacctactg gattaattta tggcaataaa atttttttt gtcttttca 420
qttttatc
                                                                  428
<210> 139
<211> 160
<212> DNA
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<213> Homo sapiens

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<220>
<223> Genbank Accession No. AA328993
<220>
<221> unsure
<222> (1)..(160)
<223> n = a or c or g or t
<400> 139
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tactaaaatg tgaccctcat tttnctttac atgaaagaac atagaatatt tcacaatgca 120
tcccacgtgg taagaataaa aaattgtttt agttatatgt
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<210> 140
<211> 359
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA342337
<220>
<221> unsure
<222> (1)..(359)
<223> n = a or c or g or t
<400> 140
agagataacc agtttatttt ggggagcaaa gagaaagggt ccctaacccc agactgcctg 60
cgaagaggtg aaatggaatt gaatgggatt atggtcagcc aaggcttcct agtggagctg 120
ctacctganc tgagttttaa gaggggtagg aaagaaaaaa tgtagtgggt cataatggca 180
ttccagatac aggggacaca aacagctctg tgtttatgaa ctacaaccag ttgttgactt 240
ttgtttcaag tggctcccct tccccagtgc tgtgtggacg atggactgaa gaggagaagg 300
ctgggagcaa gggaccagta agctgttgca gcagtgcagg tgagatatga ggcctcaac 359
<210> 141
<211> 346
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. AA347359
<220>
<221> unsure
<222> (1)..(346)
<223> n = a or c or g or t
<400> 141
gtgttgcaaa gcctttaatt agaatgtttg tattttttac atcatgcata acttcacatt 60
tgtgattaat tagtaattat ttcaatactt gtaagcncat ctgcctcaga tttaatcata 120
atacatgaat taaattaatc aaattaagga acagcaattt agaaagaaac acactttaag 180
aaatcaaaat totcaattca ggcagtotgt ttotatcatt tggtattcta ctcctttaaa 240
aatttcatat tgcccaacaa aaagtggtta tttttactgt ttttggagat gactgaacag 300
atgaagggca tcagatgcct tcatcagctg ggtattttqc ctaaga
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<210> 142
<211> 196
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA350265
<220>
<221> unsure
<222> (1)..(196)
\langle 223 \rangle n = a or c or g or t
<400> 142
caatagcaga cttttaatca atgccagaga caaagtgagg ccgagctaag aacacgctca 60
gctncgttac aatgaagaaa tggtttcctt tcgatgcaaa gtataattgt aaaccacagt 120
gctcgcacag ttcacgnctg nttaaagnga aatcttagcc atacatcacc taaaagtaat 180
taaaaagtca acacag
                                                                    196
<210> 143
<211> 286
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA358038
<220>
<221> unsure
<222> (1)..(286)
<223> n = a or c or g or t
<400> 143
caggttattt ctctttctcc tttttaatgt agagctgcag atacacttaa gttgccatag 60
taatggcaga aggagggaag ggtgttttct ttgtaaaatc attggngtat acaggatggc 120
ttggcaggta acaacactat ttctacgata tctacttatt aatataattt tatgttaata 180
tcccattctc ctcaccataa tcaccataat gttcaaattt taattttgta ttcattttga 240
atgtttgcat gtgaaaaccc aactaatcta ttatttcaac attaag
                                                                    286
<210> 144
<211> 287
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. AA374109
<220>
<221> unsure
<222> (1)..(287)
<223> n = a or c or g or t
<400> 144
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ctecteettt cecaacettg ettettaggg geeeegtgt eeggtetget eteageetee 120
tectectgea ggataaagte ateceeaagg etecagetae tetaaattat gteteettat 180
```

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aagttattgc tgctccagga gattgtcctt catcgtccag gggcctggnt cccacgtggt 240
tgcaqatacc tcagacctgg tgctctaggc tgtgctgagc ccactct
<210> 145
<211> 292
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA380393
<220>
<221> unsure
<222> (1)..(292)
<223> n = a or c or g or t
<400> 145
catggagtca gggacatgtt taattcattt gtgaatcccc tggtactggc acatagaaag 60
cgtcccatat tatctgcaaa atgaatgant gaataaatga gcaagtaggt gaatgantga 120
ttctnaggtc tcctccagct ttgatggcct atgaccgtgt gactcctgca tatgcatgan 180
cacacagaca cagacactac acacatgcac agacacacat acacacttgg ngcaaagagg 240
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<210> 146
<211> 255
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA382275
<220>
<221> unsure
<222> (1)..(246)
\langle 223 \rangle n = a or c or g or t
<400> 146
aaataataaa tgaaagattt tattcatctt tgtagataac aagcactcaa aggttaatga 60
gtgaaggaga taaccatctc ctccaaacaa agnggctctt aataacgcag aagcaaaaat 120
ctttccactt ttagatgaaa acaaactaaa aaataacttc aggcttcaga tatqqaaata 180
aagcaccatt tttcaaatgg tagacttggc ttacttaaaa taagtaaata gcccccgnct 240
atctgaaaaa gaaaa
                                                                     255
<210> 147
<211> 407
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA386264
<220>
<221> unsure
<222> (1)..(407)
\langle 223 \rangle n = a or c or g or t
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<400> 147
ttatttaata actgtagaaa tccaaaagaa ttagcatcaa atcttgaagt cgtgagtnaa 60
gctgcgggtt ggcttgactg ggctcagcca ctgagctgcc tcaaccggcc aaggaacggg 120
attatgatga ctatgcggac ttctatattg tcttcatctc attgtgtgta ttatgtattt 180
agtttcaata aagcatttgt accaatggct ctggagcttg gaggaagact aaaggaatgt 240
gtagtgattc tgaagtaaga tgtagaccta cgcagcagag ctatggggga gaagattaac 300
aaagtccttt cttccaatat caggatagtc atgagttgca gtcccatcca aaaggtcatt 360
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<210> 148
<211> 205
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. AA386386
<220>
<221> unsure
<222> (1)..(205)
\langle 223 \rangle n = a or c or g or t
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cctganganc tgtnagcgtc tgattcagct ccagcatcct tcttcaggcc aaagaactcg 120
aggatgcgct ggttgtcggt gtggtcgctg tcgatgaaga tqaacaggat cttgcccttg 180
aagctctcgg ctgctgtttt gaagt
                                                                    205
<210> 149
<211> 440
<212> DNA
<213> Homo sapiens
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<223> Genbank Accession No. AA397919
<400> 149
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tttgtgagat ggcacatagg caggtttggt gtttcctaac actatgaata tcttaaattg 120
cttttgaaag ttttatccac aaagaaagaa aaataagggt ttcctcacag ttgaaaatag 180
tttttgaaaa aaggttaaga ggaaaaaaat ctaaatacca tccttgataa agaaatggaa 240
cttcaaqtta aaaatacaaa tttaaatgaa qttttataaa atattaaaaa ctaqctaaaa 300
gtacatgcat aggcatttaa tcaaggtaag aggaacagca gtggaactta aatatgatac 360
aatttatcaa caataaataa acatttcagt gcaaatagtg cagaaaaatt tctcaaagat 420
catagcaatc attctaatcg
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<210> 150
<211> 425
<212> DNA
<213> Homo sapiens
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<400> 150
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cagagggaac caccagagaa acagcatttc agaattgtct ttcctttggt gtatggatat 240
gtgtgtgttc tagtctttgg tgggcaatgg aatctgcagc tccatgacaa tcttgttaag 300
tagcttatgt gggaagtgtt tcaggtcaca agggccaccc attctaaggc ttctcactta 360
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<210> 151
<211> 382
<212> DNA
<213> Homo sapiens
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<400> 151
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cccggctcca aagaacattt cttaagattg gtggtgcaag gatcacacct tgagaaacac 180
tgatttaggc cttcccacag tacaaagaaa tgttgcctgc cccatcctta cagcacacct 240
gatgacttac aagaggtgct gctgaattcc tcccagggaa gcaaccttaa ttcttctcag 300
caagacaagg aggcagcctt caggaaggac ccaggagctt ggtattagag gatgatccaa 360
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<211> 449
<212> DNA
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<211> 295
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<400> 311
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<400> 312
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gccctcctct gccccatgcc tttggggtct gttcgtccgt cttttttgtt gttgttttta 180
tatattgaag cgcctggccc agccccagc ccccagccc cgcactgsgg ttaatttatg 240
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<211> 1425
<212> DNA
<213> Homo sapiens
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<400> 313
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<211> 493
<212> DNA
<213> Homo sapiens
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<220>
<221> unsure
<222> (1)..(493)
<223> n = a or c or g or t
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ncctttgcgt attgtggtca ctcattttgt tgtacaggaa cagctttaan aactatgtca 360
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tcactcccan aatcttctgc aatgtgtaga aaagcagcaa ggactagatt gcctagggga 420

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<211> 3198
<212> DNA
<213> Homo sapiens
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<400> 315
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cagattgacg tcttaacaga gttagaactt ggggagtcca cgaccggagt gcgtcaggtc 240
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gcatccactg ctacagctga acagtttttt cagaagctga gaaataaaca tgaatttact 360
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gacaatggga tgttttcacc aagtggagaa tcgtgtgaag atattgatga gtgtgggacc 1920
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gaatgttctt tcattaaaag accaaaaaga agttaaaact taaattgggt gatttgtggg 2640
Caqctaaatq caqctttqtt aataqctqaq tgaactttca attatgaaat ttgtggagct 2700
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cacttaatat tatttggttg aatttgttca gtataagctc gttcttgtgc aaaattaaat 3180
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<210> 316
<211> 217
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. F01920
<220>
<221> unsure
<222> (1)..(217)
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tccaagtcag taatggaaat atgcaagang ttcaatttag gtgaggtgaa tttttgcatg 120
tgctttaacg gttgaggttt agtgtatatt gtacttttta cccttaaggc caagtaattg 180
gcaactgtga accattaatg taaaatattg ataataa
                                                                   217
<210> 317
<211> 205
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. F02204
<400> 317
caggagaagc ctgtttatta ggcaggagaa gcagcagggc agccaggctc ccctcccagc 60
caccagctgg ccaaatgtcc tcccttaact caggggtacc caaggetcca tggccatgtg 120
accagaggcg tgtaccctca agaggcggcc cctcagccct gggcagccca gccactgggt 180
ctcgcccttc aggggcctgc gcccc
                                                                   205
<210> 318
<211> 298
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. F02245
<400> 318
gggggtggca gtgcacttta ttaacaaaca aaacagtacc atacaggcaa aatcttactt 60
cagtggcaaa gcacacacat aggtatactc caacgtgtag cactggggca aacttcagac 120
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atggaacatt aggcaccaag ttcacaatca cactaaacat agttcacaat ccttcaatcc 180
atactettea qtqqaqqatq aqqeettatt taacagttaa ctgggacaga cagatgaagt 240
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<210> 319
<211> 212
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. F02333
<400> 319
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gcacaaggcc ttgatttcat catgcttttg ctgtggatgt agtgtagctt gctgaacagg 120
tatggaagct gtctttgctg ttaagtactt ctcccgtttg tttatcaacc tgcagctaac 180
aggatgtctg cttttttaca ggtttatttc ac
                                                                   212
<210> 320
<211> 221
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. F02470
<400> 320
qtttcacatq aqtqaaaaaa ttaacaqctg ccctcatttc tgaaaacaaa aaactataaa 60
caatcactgt tgctcccaat gggaccgttg gacataagcc ctgaggcttt ggggtcaacg 120
ggctagactc tagaagccca ggaccccgcc aaggtcatgt ctgcatactt ggggcagggc 180
gagctgttga accatcgcat ttctctgctg cttctttaca t
                                                                   221
<210> 321
<211> 312
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. F02992
<220>
<221> unsure
<222> (1)..(312)
<223> n = a or c or g or t
<400> 321
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gaaaaaaaacc cccgccaaat ctgaaccgcg ttgtagctcg gtccccgcct cctcagcggg 120
ctgtcgcgtg caacaaacct ccccatcat cttagaaaat aattatagag cgcggcgcc 180
cgccctcgnt cctgccagtg ggcgntttgg tcctattttt tggattattt cattacgaag 240
cacgtgaatg aatctagccc ccacaccttc aagaaagaaa ctcgcggact ggggttgaaa 300
agcccaggtg gg
                                                                   312
<210> 322
<211> 202
<212> DNA
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<213> Homo sapiens
<220>
<223> Genbank Accession No. F03254
<220>
<221> unsure
<222> (1)..(202)
\langle 223 \rangle n = a or c or g or t
<400> 322
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aaatagattt taactaaaaa attatttcgn gacaaaaata acaatatatg tnaataaaag 120
gctcaattaa aaatgtataa caattataaa cacatacaca tcaaacaaca gtnccccaaa 180
atacataaag caaacattga ca
<210> 323
<211> 305
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. F03969
<220>
<221> unsure
<222> (1)..(305)
<223> n = a or c or g or t
<400> 323
gaactttggg aaaattattt atttctcccc acggggttca gacaagtaat ttcacatttc 60
attgtaagtc aagggtaaga aaacattttt tgtacatcca tcactaatag agatcacagt 120
atgtcaatga aatatttaaa tacactgtac agagattgct ttttaatgga tttctataag 180
tagtattaat aggaaaaagc atataataca atctactctg tatctaagag ctttaattta 240
ttcaaatatt ggaagaaatt catctnctga attttnctta tttaaaaagc attatgagaa 300
ctgat
<210> 324
<211> 335
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. F04112
<400> 324
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gctgcagcct ccacctcctg ggatcaaccc ctacctcatt ctcctgactg ggactacagg 120
cactcaccac cacactgggc taattaaaaa aaaaaattct tttttgtagg gaagtggtct 180
tgctatgtca cccaggttga tctagaactc ctgacctcaa gtcacccgtc cgcattatcc 240
tcccaaagtg ctgagattac agacgtgagc cactgcactt ggcctattta gggcttctaa 300
ttcactttcc ttttccttct tgtctaattc ttgtg
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<210> 325
<211> 178
<212> DNA
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<213> Homo sapiens
<220>
<223> Genbank Accession No. F04492
<400> 325
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ctgccgcatt qqcctctcaa agtqctgcga ttacaggtgt gagccattgt gcctggccaa 120
aatgtgtatt tttaatatgc tgctgagttg actcttgtat gatcaggagg agcatttg
<210> 326
<211> 211
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. F04816
<220>
<221> unsure
<222> (1)..(211)
<223> n = a or c or g or t
<400> 326
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aacaattggg taatttgtga gacaccaaag aaaaaaagaa tgcacctatg agttacagag 120
tccaaactga tcagggctga caacttgacc accatgtntc ccacaccacc acccccacca 180
                                                                    211
ccaccaccac caacagcttc gtcctcagag a
<210> 327
<211> 276
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. F09281
<220>
<221> unsure
<222> (1)..(276)
\langle 223 \rangle n = a or c or g or t
<400> 327
actgtttaaa tataattgaa gtttttnata tgatgaagtg ctccataatt taaatgtaaa 60
aaaccaatag gaaatatatg aaataaaata aaattatacg taaaagtgac aatgcctcta 120
ttagatttaa cagtatctta caatagaata agttgaaacc tacaaaatgg aagaaagttt 180
aaaattaggc agatattatc ancctggtga agaataaata catatgtcaa taagcattta 240
atgtatttgg tcttagattt tacatgaaat aataaa
<210> 328
<211> 293
<212> DNA
<213> Homo sapiens
<220>
<223 > Genbank Accession No. F09315
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<400> 328
acagaaattg acctttattt gttgtactaa agcctgttta acttttgata caaagtaaca 60
ttttagtaca gaaaatccca gtctgtcagc tcagtacctg tctgtgcaca ctgtaccatc 120
tcagtcccac tctgcctgta acttagaaaa cagcccctac ccccagaggt ctgcgagtta 180
ataccttgag aatagtctac agtttttcat agtttgtctg agctagaaaa cttgtacctg 240
taaaacaaag gacagcattg aggactgaaa cttgtctctt ttttgaacaa ctg
<210> 329
<211> 214
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. F09684
gctttacata aacttataag gattttttat ttaaaggatt taaaaatata acacagtcaa 60
tataaacatg tactgggaat tataaaccat tctttcttct aagcactgga tgagatacta 120
aaaacataca gtatcttacc aatagccatt aaaataggct aaaatgaaaa agaaaccgtt 180
gtaacaaggt tactaatccc ccaactttca atgc
<210> 330
<211> 332
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. F09748
<400> 330
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tcaactttta aactcagcac tctgttggag tggaggtgca cggtccttca tcataggcag 120
cctatgcgag atgcatctta ggaagggagc tttcgctgct cagaaatcaa agctccatcg 180
aaggggaaag ttgacaacaa ttcaggggct ttgagtagtc aagacaatta gcttagtact 300
tcaggtcaat aaatgctaca atttatgggc aa
<210> 331
<211> 247
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. F10078
<220>
<221> unsure
<222> (1)..(247)
\langle 223 \rangle n = a or c or g or t
<400> 331
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ttcagaagca ggaaatttta gttgttgcca gagaggtgtg tcaaggacac agtgaaagga 120
gccatgcgga catggggtgg aaggetttnt ccaacactgt tacaacactt ttgtaaatga 180
gcaaaacatc tttaaaaatc cttataaatt ctttataata tgttacacat ttagagacaa 240
```

```
tatttac
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<210> 332
<211> 243
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. F13763
<400> 332
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caatcaagat agtgtattat tagaaataac attaatagaa gcttggtcag aaatgataat 120
agtcataata agcatctctc tcaccaaggc attccacaca gagagatcac agcacaataa 180
ataaaggatt tctcatttgc cacacaacaa ataaaacaat tgcagtaaca aaaatatgac 240
ttt
<210> 333
<211> 415
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H01824
<220>
<221> unsure
<222> (1)..(415)
<223> n = a or c or g or t
<400> 333
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gcatacattt tttacagaca atatataaat gttgtacata attaacaata acttagttca 120
ctaatccaaa ataaaacaag ccaaataaaa cataaaaaca gaaaatactg ccgnttcttt 180
ttcttatgcg ggacactagn tacaaaataa gttacttctg ggccgtgggt gctccctgca 240
ggegactgcc cgcccatatt gcacttgggt cactaacatc aggcacaatc ctcctccggg 300
ggccggggcc ccttcancag ggcccaccac accccgccgt tcaccggcat tacaggaatc 360
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<210> 334
<211> 309
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H02308
<220>
<221> unsure
<222> (1)..(309)
<223> n = a or c or g or t
<400> 334
tgatagcaca ttttagtttt taataaaatc tgctttttac ttatatttaa ataaattgcc 60
cagttactga atcagaagca tttcttacaa agcaaacaaa ataagcatcc cttctatgtt 120
aataacatgt taatagtatg ttggcaagtt gatttagaac aacttgccaa caatacaaac 180
```

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agaaaaaagg agtgggtcaa agaaatctag tttggcttta ttttcaatag atcatactgt 240
ctgttgaaaa aggaataaat aattatggag cctatctaat aatatactca atagnttgaa 300
attattgag
<210> 335
<211> 277
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H03387
<220>
<221> unsure
<222> (1)..(277)
<223> n = a or c or g or t
<400> 335
acgcaagtta gannanttat tatgataact ctgcaatctt ttcagccact ctttaaggtt 60
cctgggcatc cattctgggc acagtgtgac atttacctga acagagagga gantggcact 120
agaagatgag ggagatttgg tgcctaaaaa ttactacaaa caggcagggt gcagtggctc 180
acgcatgtaa tcccagcact ttgggaggcc gaggtgggtg catcacgagg tcaggagttt 240
gagatctgcc tggccaacat ggtgaaaccc catctct
                                                                    277
<210> 336
<211> 372
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H05084
<220>
<221> unsure
<222> (1)..(372)
\langle 223 \rangle n = a or c or g or t
<400> 336
tttttttttt ttcacagtga gcattaaatt attattccat acagccctgg ccctggccct 60
tcttgaggga gtggggtttn tggggtntgc ccagcaggga tcctgccaga tgatgtccac 120
atgagaaggc aggtgtccaa cagcttcagc ttcacccagt gccccccaga caaataatga 180
caagtccaqg gtcttctgat gtgtcaggcc agcactcccc ttgctgatgg gaaaaccggq 240
gctcggccag ccccactgca tcccctcaca tgatgatacg aggctctngc actgactcgc 300
caatagactt gtggggcagc angctggctc cgttgaggta ggagctcatc attaactatt 360
gacgtcctnc ac
                                                                    372
<210> 337
<211> 353
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H05625
<220>
<221> unsure
```

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<222> (1)..(353)
^{\circ} <223> n = a or c or g or t
 <400> 337
 tttttttttt tttttttt gcttcacaaa tgtcaatttt attgacacta gtgcacaact 60
 aaatacaata attgcaaagg aagtggaacg tgttcaaaca gaaatggtga caatgagtta 120
 qaactgcagt tntttcaaqq tactacacta ttatttaaaa aaaaaatcac aaanaqaaaa 180
 atgttatcac tacaaqtaqq gatttaqqaa qnqaqnaaat tctqqqcaqt ctqtctaqna 240
 gggttaaaac atttcatggc atttgtgagt tgctgttgga gagttgtttt ttatttgtcc 300
 accgtaatct gggcaacatc cgggggctta ccttcagctc tcggcactgt gcg
 <210> 338
 <211> 501
 <212> DNA
 <213> Homo sapiens
 <220>
 <223> Genbank Accession No. H05704
 <220>
 <221> unsure
 <222> (1)..(501)
 \langle 223 \rangle n = a or c or q or t
 <400> 338
 tttttttttc cttctgtagt cgtctttatt tagagcagaa ttcagactca gctggtatcc 60
 cccagggcaa ccccaggatg ggganagggc tggtctgtcc ccacccactt ctccaggatc 120
 ctcccagccc ccaggctgnc ttttccctcc aactgtcagc tgcttagctg ctcatctggg 180
 gattggagct ggagcatctg tcaaggttgt ctccttgaca aacagcttcc tctttggaaa 240
 tggcttcact caggtcctgc aggtcatcga gcaggacaga gagggacccg gggaaggaag 300
 acagcagatg agcaccagac aagggaaggt gctcgtggtt acagagggaa acagggttgg 360
 gcacagggaa atgagggaat ggggagagag ggaggctctt tgggtccaag ctggggcatc 420
 ncttaaaaga ggtttaaggg tntcgaagga ccncagagaa caacattctt cntgcgagat 480
 ttttaagagg gagttttctn a
                                                                    501
 <210> 339
 <211> 465
 <212> DNA
 <213> Homo sapiens
 <220>
 <223> Genbank Accession No. H08548
 <220>
 <221> unsure
 <222> (1)..(465)
 <223> n = a or c or g or t
 <400> 339
 ttttttttca caaatattgg cttggttttt atttctatgc ttataaaaaa aatatgaagc 60
 ttctttgtgt ggactgaagg ggtgttagcc tgtggatgtt ggtcttcggt gcctgtaccc 120
 cagtggctgt ttacattcca ggnccctgct aaataaagna ggctccactg ccagctgtct 180
 gtacactttt tcttggggga agagttcttg tcttcagttt actgcagtag ggttcctggc 240
 tctgttacat gctcatgtgt tccggaagaa catatgaaat atcatcccac ggatgacgat 300
 acagecectg cttcagectn ttctgatcaa gatagtntcc aatqaacecc atacteettc 360
 ccagcacaaa gatgccattg agggctccaa tgtcaatatt attgcatcag cttcctcccg 420
```

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agtaaaggga cccacagttt tttaaggatg ttttacaatt gcgat
                                                                   465
<210> 340
<211> 313
<212> DNA
<213> Homo sapiens
<2220>
<223> Genbank Accession No. H15143
<400> 340
tttttttttt tgtgggtcac agttgagggt ttattgccag tgttaggaag aatggggggt 60
ctgggtggcc aggggtcttg ggaggaattc caaatgagca ctgcagggcc tgtgagtggg 120
gaggagaget getgececec tgccacccag gaggececag ggetgatgec accatatect 180
gactgctagt ggtgccttaa aaggtggcct ccccacagga ggggagcctt gggggccccc 240
aggagtcagc cetcaccaac aagccetete teaaggggge caggggettt tatteeteat 300
gggacaggct ggg
                                                                   313
<210> 341
<211> 295
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H16171
<220>
<221> unsure
<222> (1)..(295)
\langle 223 \rangle n = a or c or g or t
<400> 341
tttttttttt ttttttaaa ttaaacaccc ntatganttt attaaatcca gaactgtgtt 60
aaagggcggc ggtctncgag ggggagtntg gtagggggac gagggacaag atgatgaacg 120
geogtgggea tecentaggg ngacceggne caccecegee caacceacce cetengeaac 180
gctgcatcag cttcaccatg attcccagtg gtgctgggct gggcagggcg agatggctgg 240
gaaacacaga gggacagagg gacagacaga cgccttccac aaacaaaccc tggnc
<210> 342
<211> 389
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H16676
<400> 342
tttttttttta gttttgtggt actacatatg ttttattaaa aattcaaact ttttttcaga 60
tcgaagcata atttatcttc cattaacaaa aacgaagatc ttaaatttga cacgattaca 120
attaaaatgc tgaaaggagt tatgaggcat ttaaatcatt cttcaattag aatgtttgca 180
gcatatttct cagaggctga cctggaacac attacctttg ttggcaggca tcaaaggcag 240
gataaatcct gtggctggaa atcaattgtg agtcccatta ggatgacttt ctaggcacac 300
atgcataggg tettgcactg tatecgttet acttetagga aggttgctgt etggaagget 360
ctttcccctg ggcgaggtca ctttcccgg
                                                                   389
```

<210> 343

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<211> 471
  <212> DNA
  <213> Homo sapiens
  <220>
  <223> Genbank Accession No. H16768
 <220>
 <221> unsure
 <222> (1)..(471)
 \langle 223 \rangle n = a or c or g or t
 <400> 343
 ttttttttta atttataaaa atgaaaagtt tatttgtctc atggttctga caggctgtac 60
 aagaaacatg gcaccaacat ctatttctgg tgagggcttt aggctgcttc cactcatggt 120
 agaaggcaaa aaggagctgg catgtgcaga gatcacgtag ncaagagagg atacaaggag 180
 atttccaggn ctctttttaa cagtcagctc tcatgagaag taatagagga agnaagtcac 240
 ttactactga gagagtggct ccaagccatt ncataaggaa tcaaccacca tgacacacta 300
 gggcctcacc tccaaaactg gggaatcaca tttcaacatg aggatttggg aagggtcaaa 360
 tatccaaact ataggcattc tacccctgga acgcctaagt atcctgtcct tctcacaagg 420
 caaattacat tattttattc ccattagttt cccgaaaact taacttgttt t
 <210> 344
 <211> 354
  <212> DNA
  <213> Homo sapiens
  <220>
  <223> Genbank Accession No. H17333
 <220>
  <221> unsure
 <222> (1)..(354)
\cdot <223> n = a or c or g or t
 <400> 344
 ttttttttta attgttaata ttgctaattt gtacaatggt taatgatctt ataaaatagt 60
 tgtatgaaag caccaaccac cttagaaagt ctgaccagca ttcatatcta ctttccagac 120
 cctcatccct cctccccact cacctgactc tgctcggctc attcatgggc tttcctgtgc 180
 tetgecattg etcaggtgag tgagcagtte geeeggcaca ttgaccagge agatecaggg 240
 cancegateg gtggageeca ggaaatggag aggetggeac agetgeagea atgeetgnaa 300
 getgteetga tttteteegg ettngagata gecaccaett ttgageatta ttae
 <210> 345
 <211> 486
 <212> .DNA
 <213> Homo sapiens
 <220>
 <223> Genbank Accession No. H17550
 <220>
 <221> unsure
 <222> (1)..(486)
 <223> n = a or c or g or t
```

```
<400> 345
ttttttttat ttttaaaaat ctatttattt atcaaaacag tattggcaca gtaattctca 60
tattatcatc aaataataaa attgctactt tctgtactca attctttaga atcctagaaa 120
ttgcaaatgc attcaattta acaatattgt aaataacaat acaaaagaaa gaactctgca 180
tatttatgga aacattgttg atggtacagt tctactgaaa ctcatacaca tttcactatt 240
taatttacat atggncttgt tgaaaaaaac cagtatgttt tactttttca atttccttat 300
ggctaaaata catgtaattc taaagggata tctcttgggt gttataaaaa ccagggaggg 360
tccaccacca ggtcaaggtt ggngtcaagg ntacttcaaa ggttccctqg aatggatccg 420
gaaaacaaat tttaacccna aaatgtggta ccgntttggg ggggcccttc ncgggccccc 480
caacgg
                                                                    486
<210> 346
<211> 371
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H18947
<220>
<221> unsure
<222> (1)..(371)
<223> n = a or c or q or t
<400> 346
tttttttttt cttttttag gnttcatgtt tgttttattt aaagtctggt tgggtacaga 60
aaacacacac acacttaaca ggttaaaata tccaaataaa atttactgca acttttqtaq 120
aattttattt gtgctacaag acacgttgca taagaaacta tttaaaqccc ctqaqqaaaa 180
aatatccatg gtttaaggtg caactggttt tgtttcttct ttggggaaaa ggtgatagat 240
ggtctctggg agaaattatg gggtggagtt gagaagcaca atcgaaggtt atatggtggg 300
atgattggcg aattgtgtgt cctgggttct tggcagcatt aaaatagcct aatgttttgt 360
tcttttttc a
                                                                    371
<210> 347
<211> 187
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H21814
<220>
<221> unsure
<222> (1)..(187)
\langle 223 \rangle n = a or c or g or t
<400> 347
ttattgaggg tttattgagt gcagggagaa gggtcttgat gccttggggt gggaggagag 60
accectecce gggatectge agtetetagt etceegtggt ggggggtgag ggatgagaac 120
ccatgaacat totgtagggg ccactntott otocacggtg ctccottcat gtogtgacct 180
gggcagc
                                                                    187
<210> 348
<211> 432
<212> DNA
<213> Homo sapiens
```

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<220>
<223> Genbank Accession No. H22453
<220>
<221> unsure
<222> (1)..(432)
\langle 223 \rangle n = a or c or q or t
<400> 348
ttetettgtt getggagttg taaaaateaa tgteeeattg etgagatega ageteeetgt 60
gtctctgggg ggctcagcag ggacgatggc ctccagagtg gacctctgag aaattgcaga 120
ggcatcagag ctgtgggctc agcatatgag gtccccaggg gccatagacc ccctcctct 180
gggaagagtg ctcctgcaga gcttatttgc aatctcctgg gagtcccaga ctcaccaaag 240
qattcaqatc ctcttcttt tgcctcctac atagagcaca ttatagacct qaaacaggaa 300
tcagaattcc agactccctt agtgaggaga caaagtgtta ggtcttagct ttttcccttc 360
taaattaagg gtcctccctg ggattcaggt tgcctgatag cttatncctg aaantggtng 420
gagataggga aa
<210> 349
<211> 233
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H26288
<400> 349
aaaaacacca gtttgaaaca cattactgaa agtgagtgta cacaataaat agaaaatagg 60
gatgcatagt gctggagaca ttcaaccaac ttatcttcat ctgttgccta ctgttgtaga 120
caaaatttga cacacaatta gcattactga aagagcagcc aaactacctc ggagaaagtg 180
ggcaaactac tggaaaagta gcttaaagct ctgggaccac tcaccaaaaa taa
<210> 350
<211> 290
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H27180
<220>
<221> unsure
<222> (1)..(290)
\langle 223 \rangle n = a or c or g or t
<400> 350
aggntttatt ttggaccaaa aaaaaaacca caattgtttt ctagctggaa gantgggcaa 60
ggggggtccc agacagtaaa ctcccccacg ggtgggttga gcctcaggtg gggggtctcc 120
tgttgtctgt gcttccccac acagcagcct ccctcctggn gtctgtggca gccacgggag 180
gggcagacta ggaggagctg ccacagttnt tcacttgggc aggaagtcag aggactcaga 240
caccagette ceategeggg thtegatett etthanaace acggeeetgg
<210> 351
<211> 292
<212> DNA
```

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<213> Homo sapiens
<220>
<223> Genbank Accession No. H27675
<220>
<221> unsure
<222> (1)..(292)
<223> n = a or c or g or t
<400> 351
gtgtctccat ggcgagtggg agcgtgaaga tgaccagctt tgcggagagg aagctccaga 60
gactcaacag ctgtgagacc aagtccagca ccagcagctc ccagaagacc acgccagatg 120
cgtctgagag ctgcccagcc cctctgacga cgtggaggca gaagagggag cagagtccga 180
gccagcatgg caaaggntcc cgccagcctc ctggcatctg agctggtaca gtggcacatg 240
cantcgaagg agaagcgcag ggccatcgag gccaggaaga agaagatgga gg
<210> 352
<211> 327
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H40424
<220>
<221> unsure
<222> (1)..(327)
\langle 223 \rangle n = a or c or q or t
<400> 352
ctgtatantt tnncttnttt tttctcttgt gatttggcac ttaaggctta agcgcnaaaa 60
aaaaaggcat ctactgacaa aatatgggac ttgtctgtna tgcatggtaa gtgggctata 120
aaatccaggg agggggtttc aagccagaag aagctactga caaattgact tgtccttatg 180
ttaggtgggg ttatgagggg gagagggagg gcacattctg aggtgctggg ggaaaggggt 240
tgagcttaac cttgttaatg tagggcctgt ggggaatggg atgggtaggg agaagagggt 300
atgggatgtg ggtgcagggt aggggct
                                                                    327
<210> 353
<211> 448
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H44631
<220>
<221> unsure
<222> (1)..(448)
\langle 223 \rangle n = a or c or g or t
<400> 353
actcagcatn cnttttattt tnctatctga catttctaac aaaacgccag ggaqacggaq 60
ttaaaaagaa tccaccccac gaaaggtaaa caaaggagac cctcagaaac tccctggcaa 120
ggatgttccc ctccccagat tgggcccagt ttcaccagca actgggtctc agactcagcc 180
ttatgccttt ccactgacac cccccacccc tccacantct cgtgattcag accagggaac 240
```

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ttctcgggct gattgtgtcc gtgtgtctga gggagggca cgctggaacc tgggaaccta 300
Ctgggcacct ctaatgcaga tgagaaaaac ttgagaatgt gaaaggagat cagtccccgn 360
tcccacccga aggtgcagag acgcgggaca ttaaccagca gnacgcgggg gtgaaggaac 420
tcagggcaat ttctcccant gccagggg
<210> 354
<211> 346
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H48793
<220>
<221> unsure
<222> (1)..(346)
<223> n = a or c or g or t
<400> 354
gatttaggag attccaagtg atacctttaa ttcactactc tatgtcctta ttaataaata 60
catatttaaa aaaacctata caatatagtg tatttacagc atggaagagc agagactctg 120
aagccagact gcctgagttc aaatcctgac acttctactc aaatatgtgt gagtgacttt 180
gggcaattta cttactcttt ctgtgtttct atttactcgt ctacaacaat aatttctacc 240
tcatcaaatt aaattaaaaa aaaaacggct taaataggtt aacatttgta aataggctta 300
ggaaaacact acatttaaaa aaataancat tcctaaccca ccttcc
<210> 355
<211> 458
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H49440
<220>
<221> unsure
<222> (1)..(458)
<223> n = a or c or g or t
<400> 355
ggagtttcac catgttggcc aggctggtct caaactcctg acctcaggtg atccacctgc 60
ctcagcctcc caaagtgctg ggattacagg catgagtcat tgctcccagc cattagaaaq 120
attgttaatc ctatgaactc ccttttgtag gagagaaagg gccaatctgt aggggtagcc 180
ctgtccaggt aaagttgttt tcagcctcat gtctactgtt aggtgaggga qtcacagcca 240
gacagagagt attgctggag ggtgagagaa ttgtgqagac caactaccac ataqcaaqaq 300
cccagctctt gggagcattg agatgtaagc tcagggttac acagttccaa atcttgggga 360
aggggctttt tcagacagac tgtttgcttt ctgctgagat taaggaattg catcantctg 420
ccagagtatt gactttttaa cagattatta aataaagg
<210> 356
<211> 446
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H52835
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<220>
<221> unsure
<222> (1)..(446)
<223> n = a or c or g or t
<400> 356
cgqataccct gggggcctct qctcctctct ttqtqqaqac qtcqtttcac cgqcqqcqcq 60
tgaccccggc agctgtccag agacccagag atgtccaatc acaggcgcac ggtgcacagg 120
cgcgcagggc tgcctggaac gggcccaggc aggcagtgac cgggacctct ccggagggag 180
aggaacggtg ccctcccggg aggagctggc caggcaggcg ctgcccaggg cggccttccc 240
tgctggacta cggcattgcn actgagttat ataaagacac tatttgggga aggacagcgg 300
gtgaggactn ggcgcggcgg cacacgcttt gcctgttgtn ttcagctctt ctgggggcca 360
aggcagggag ttccagggtt tacagtgagc ctgatngcca attgctttcc aaaagagaga 420
aacagagaga aagggattna ggcttc
<210> 357
<211> 386
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H54764
<220>
<221> unsure
<222> (1)..(386)
\langle 223 \rangle n = a or c or g or t
<400> 357
gatggagttt cgctcttctt gcccaggctg gagtgcaatg gtgcaatctc ggctcactgc 60
aacctccacc teetgagttt gagattetee tgeetcagee teecactggg attacaggeg 120
cctgccacca cgcccagcta attattgcat ttttagtaga gatggggttt caccatgaaa 180
atttttattt ttattaaaag agtgcatgag ttagtcatga aggcagagcc agggcgcct 240
gcataccaaa tgtgaaggaa cagtaccaat tgacaaagga aggcacaaaa ctaggacaaa 300
ggaaaaggga cttcaattaa ataaggtaat ttggaactaa ctggaaaatt gaggagggg 360
aaatngcaaa taaaatnggg gaggca
                                                                   386
<210> 358
<211> 384
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H56673
<220>
<221> unsure
<222> (1)..(384)
<223> n = a or c or g or t
gttaccaaga cacaatttta agatcaaaca agtgtcaagg taggccatgg cttgttggca 60
gtagtagggg coctatggct atttccaggt atgggtggcc ccttttcctt ggttatctgg 120
ggaatctgcc acagcagaca gcaaaaggta aaaagcatcc ctttaataac tacaccccac 180
tccagcaatt gaggtttatt caggggtggg tcaaagtagt acaagacaaa aatagcttag 240
```

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tgaaatggnt tagaatccag actgaggtgc cagactgcct gcatctgagg tctcaggtcc 300
caccatgtat ggaggccgtg tggaccttgg gggtgaggtt actaggcctc cccggggttt 360
caaatcttct tcacctgtaa aatg
<210> 359
<211> 440
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H58873
<220>
<221> unsure
<222> (1)..(440)
\langle 223 \rangle n = a or c or g or t
<400> 359
actataactt agtgtctgta tttaatattg acaaccaaaa atatatatan ttttnttgca 60
tctatacaca acaggcagg agtctccatg tnttcttgag cagtgagttt gcaggctccc 120
acaggccctc ttctcatggt aatagtgtgg ccctagtgca aaggagacta gaacccggca 180
gcccagactg gcccttcccc tctcctccct gcactccagt gcttcccaac tggtctcagg 240
taaagaaagn ttantttgag tggttgggta ggaagagatg ggaaggggca aatcctaatg 300
ggagcctgac ccctagagtg gggagttcca gggccagcag aacgggtggg ccatagccct 360
ncctggggnt agaagetttg tagtteatag ttegattagt ntgteentag ggeatnaggt 420
nccagcccta cagattagct
<210> 360
<211> 284
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H60595
<400> 360
aagacagagt ggactgttac aaatgatttt gcaaaataca aaaatagata tacttccact 60
gaatgcttta atcatttttc cgggcactct catcttttgg ttcttcctca tctgagtaca 120
cagtgggctc ctcccctcc ttcagcagtt tgcccacgtg atgatacttg aaagtgaact 180
gagactecca gteacteaga gteteetget gggegeagtg aggteagaaa ggteategta 240
ctcatccttc agtgcttcct tatccgggga aaatgtgggc aagg
                                                                   284
<210> 361
<211> 317
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H61295
<400> 361
quaeceteta agggacetea auggtgattg tgccaggete tgcgcctgcc ccacaccete 60
ccttaccctc ctccagacca ttcaggacac agggaaatca gggttacaaa tcttcttgat 120
ccacttctct caggatcccc tctcttccta cccttcctca ccacttccct cagtcccaac 180
tecttttecc tatttectte tecteetgte tttaaageet geetetteca ggaagaeece 240
cctattqctg ctggggctcc ccatttgctt actttgcatt tgtgcccact ctccacccct 300
```

```
gctcccctga gctgaaa
                                                                    317
<210> 362
<211> 370
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H64493
<220>
<221> unsure
<222> (1)..(370)
\langle 223 \rangle n = a or c or g or t
<400> 362
gggtgcttta tttccatgct gggcgcccgg gaagtatgta cacggggtac gtgccaagca 60
tectegegeg acceegagag eeeggggage gggngettge eggeegtege acteatttac 120
ccggagacag ggagaggctc ttctgcgtga agcggttgtg cagagcctca tgcatcacgg 180
agcatgagaa gatgttcccc tgctgccacc tgctcttgtc cacggtgagc ttgctgtaga 240
ggaagaagga gccgtcggag tncagcatgg ggaggcntgg gtnttgtagt tnttctccgg 300
ctgcccgctg ctttcccant ccacgggcga tgtcgctggg ggtagaagcc tttgaacagg 360
                                                                    370
gaagtcaggc
<210> 363
<211> 460
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. H66642
<220>
<221> unsure
<222> (1)..(460)
\langle 223 \rangle n = a or c or g or t
<400> 363
ttaaagacag agtttcgctc ttgttgccca ggctgtagtg caatggcgcg atattggctc 60
actgcaaccc ctgcctccca ggttcaagtg attctcctgc ctcaccaagt agctgtgatt 120
acaggtaccc gccaccatgg ccagctaatt ttttctattt ttagtagagc cggggtttca 180
ccatqttggc caggctggtc tcqaactcct gatctcaggt gatccacctq tcttggcctc 240
ccqtqctqqq attataqqca tqaqccacca cgtccqqcca aattttactt cttaaaaqtq 300
cttttctctc agtgatatca aggtcttctg tctactatta taaccataag cttctttagg 360
cattaaggag ggaaaatgtt taataaaatg taattaaact gggatggaat ggtcagtgta 420
tttaaatgta aatatactta aatgtaatta ccggggnggt
<210> 364
<211> 291
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H68097
<220>
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<221> unsure
<222> (1)..(291)
<223> n = a or c or g or t
<400> 364
tgaagtttat ttnctctggc agtatgtttt agtttcttgt ttttnatttt gttgtgtgtg 60
tatgtgttgt agattttatg atttgaggtt accatgaggc ttgcaaataa cataacatgt 120
tattttaaaq tqacaacttq acactqattq caaaaacaaa caqqqcqaaq aqaactaata 180
aaaactgtac actttaactt cattcctcct gtttttnaag gtttttatgg gtttctattt 240
atatctcctt gtactatttt gaaaagggna ttgcaggtta tcatttgttc a
<210> 365
<211> 317
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H77597
<220>
<221> unsure
<222> (1)..(317)
\langle 223 \rangle n = a or c or g or t
<400> 365
tcaagtctaa gtgtttaatt attattcaca tatttcacag aaaaaaagga atgtagcaaa 60
tgagtcggag ttgtagaaaa aaaaaatcct ggnttttacg tgtcattctg ttttcatctg 120
acagcaggge tgtcccqaca tcaggcacag cagctgcact tctctgacgc ccctttgcag 180
atgcagccet gggcacactt gggcacagce caggggnaaa caggagcagc agcctggggg 240
aaaaagggag agagaaggtc acaggcagac ttnaccaggg ganctccctt tcccaacagc 300
aggcctgggc tcaagct
                                                                   317
<210> 366
<211> 340
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H81070
<220>
<221> unsure
<222> (1)..(340)
<223> n = a or c or g or t
<400> 366
caggtctaaa gtgtttaatt atcactcaca tatttcacag gaaaaggaat gtagcaaatg 60
ggtcaaggtg gtataaaaaa aaaatccagg tttgtacatg tctctctgtt tacatctggg 120
agaaaggttg tcctgggcat cagtcgcagc agctgcactt ctctgacgcc cctttgcaaa 180
cacagccctg gggcacactt gctacagccc acgggnagnc agggagcagg cagctctttc 240
ttgcaggagg gtgcatttgc ctctttgcac ttgcgggaac cagcgcggtg cagggaggac 300
accagcggcg cagggagcag ttggggggtc cattngcaag
<210> 367
<211> 330
<212> DNA
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<213> Homo sapiens
<220>
<223> Genbank Accession No. H81379
<220>
<221> unsure
<222> (1)..(330)
\langle 223 \rangle n = a or c or g or t
<40.0> 367
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ggaacatcga cccataacat gcaacaaaaa tgattttgcc ttttggacat atttaacaga 120
taaacttgac attacaagta acagcaacac attcccattc tactgaagaa aacaaatgcg 180
atttaacttt caggttagaa aacgtatctt cttactgcaa tctcaagtng gcatttngaa 240
agtttagttt tcccttttct aacctctaaa agatgatatg atttttaatg caatcataca 300
caactgtttt cacattgggg aatantcacg
                                                                    330
<210> 368
<211> 419
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H81413
<220>
<221> unsure
<222> (1)..(419)
\langle 223 \rangle n = a or c or g or t
<400> 368
ngagccagaa aaggattttt tttaattcaa gtaactgaaa taggaaacca gagggggagc 60
cccaggctgg gataaatcat ggctacccct ccccaacaga acagggggag gaggtggccc 120
ctacacccat tatggtcgat tcgggccccc ttgctcactc tgctgcagca tcctagaggc 180
agggccccac cttccctggg actggggtag tcggtcaccc agcctgcatt gccccagccc 240
ctnttcccca caaagagtat cttgggggag ggnttcgtgg ggcagaacag gagggcaatg 300
agggatgaac attgctcaaa ctcctttcaa aggggcacct gaccgcacag gggaggntgg 360
gcaggaaggg caagggntgg gggatgccgt ntaaggaggg cggangcagg canttttgg 419
<210> 369
<211> 386
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H83380
<220>
<221> unsure
<222> (1)..(386)
<223> n = a or c or g or t
<400> 369
ttaattgcag aaaaatttat taaattggaa aatcttgcgt ttttcaatgg cgctggcccc 60
gggtcagcgg cgattttctc tgcatcaaga tgggctttgc gtttccgtag tgggcaccag 120
```

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tagtagecta attateagte ttetecegge atttttaagg ceagggagee gaagegetge 180
ttqtaqqcqa ataccctaca qaqcggtttq gctttttaaa ttactgttat tattttgggc 240
agagaacagt cggtctgggt gcaccccgtc ctcgctgcag aagaggctgc gagtccgagg 300
tggggtctct cgggaaggtg aaattccttc tnggggntna gcgagccccg gccccgcgcg 360
gcagtccagc ggccccggtg ttgttg
<210> 370
<211> 335
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H84761
<220>
<221> unsure
<222> (1)..(335)
\langle 223 \rangle n = a or c or g or t
<400> 370
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ngcacccatc tcgaggnggt attttcngta agancaggng ttccnccctc gtaggtttag 120
aggaaacacc ctcatagatg aaaacccccc cgagacagca gcactgcaac tgccaagcag 180
ccqqqqtaqq aqqqqccc taqqcacaqc tggqcccttq aqacaqcaqq gcttcqatgt 240
caggetegat gteaatggte tggaagegge ggetgtacet gegtagggge acacegteag 300
ggacccacca ggggactttc ttcaaagttc cnggg
<210> 371
<211> 178
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H86112
<220>
<221> unsure
<222> (1)..(178)
<223> n = a or c or g or t
<400> 371
qcttaatqqq qccaaaqqqq caacacaaaq cattqaaaac atcactqqct cacaaaacca 60
qtcaccttqt taccttctca qttqcatttq tttatttcac aaqqcttcat tcacacataa 120
aancaagata ctantccaat tcangttcat aacggttata anggtaanca tttgttgg
<210> 372
<211> 287
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H88338
<400> 372
atgcatgttt aaacatttaa tctagaactt gattacaaag taatttaatg aagaaaataa 60
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taattaaagc aattgactaa tgatctcaca gcctcaaggt tgtatgcaaa cctagattag 180
aaatactttg gtctctaaaa ataacaaaat ggaccataac atttttttc ttacaagttt 240
gaagtgggtc aattatgggg gaaacacata cattcctaag gggaaat
<210> 373
<211> 337
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H88798
<220>
<221> unsure
<222> (1)..(337)
<223> n = a or c or g or t
<400> 373
nactttaata agtataaagt atataaacaa ttaggtaagc ttgtggagaa gctgaccaag 60
atacataaat taggaaatac aagtgtccat ctaaattttc tatatttcat ttttttcata 120
atatttatta aaggtgttta atatacagtt tctcatctgt cattttggaa gtcctttatt 180
gtaaagacaa ttctattgtc tgatgacaaa cagcagccac catggttatt caggacctcc 240
acgttggata aattccattt cttcttgaga cacaagtttc cttctggtat ttctgaggta 300
atggntttta ttatttctgg cagtgtctgg tggaccc
                                                                   337
<210> 374
<211> 321
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H91703
<220>
<221> unsure
<222> (1)..(321)
<223> n = a or c or g or t
<400> 374
ccataagaca agtgacatat ccaaccaacc atccatcccc acctgtgccc tattctttcc 60
ttgtgtttct ttagagcctt ttcagctatt tcctgtgaag caaactgcac gaaggcctcc 120
cccgtactcc tcccctggaa gtccaccggc aatgttatcc catttggcac gatttccaac 180
ccttcaaccc aaggacaaat aaccccagta gggggncaat attaacatca caagcccagn 240
aaatgattct tcttataggc tttaaataaa ccaggacttt ttaactttag ggtgaatggg 300
tatgctttca acaagtactc t
                                                                   321
<210> 375
<211> 395
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. H94471
<220>
<221> unsure
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<222> (1)..(395)
\langle 223 \rangle n = a or c or g or t
<400> 375
tttgttactt ttacatgatc tttattattt aagaaaaacc tcttttaacc atttatataa 60
cagaaaaaaa atagggaggc tggtagatca tcacatatat agtagctaaa atatgaaagg 120
ccagggaatt tattattaat gaagtcataa aacagactta accaaaagtg tgtgctagga 180
aacaagcagt ttcacttcag agacttcatt gcaggaaccc agtttcctta tgtggaaaaa 240
agtgattata aataacagtt atctgaaaqg tggttgaqag gattaaatga gatcacctat 300
gcaaacaaat acatgtaggt atgaaagacc atccgtcctg ggggtngtgg aaagtttaag 360
tttccccncc agaacccttc cctttaaggg cctta
<210> 376
<211> 373
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H94475
<220>
<221> unsure
<222> (1)..(373)
\langle 223 \rangle n = a or c or g or t
<400> 376
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gageettaag gageteatgg gateetteee tgeeteggtt cetgagetee egggeagagg 120
agggagacag gagaggaagg aagggaaatg ctggcagtgt tgggatctcg aggagccgtg 180
ggaagtctgg cgtgacaagg cacaggggt aggatggagg ctgatggact ctcggcaggt 240
taggecacag ccaaggetgt gecangacac gagttecacg eggggetgag gacaacgett 300
cgcctcccga gccaccacca gggcccgtct ctccccaccc taagcctagg tgtcccggga 360
caagtccaaa ggc
                                                                    373
<210> 377
<211> 417
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H95960
<220>
<221> unsure
<222> (1)..(417)
\langle 223 \rangle n = a or c or g or t
<400> 377
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gatctaggta tgtggcccat tgcaattgag cacatttctt gggtctgttt ctctatctct 120
aagggcagtc tcaaaacccc agctcaaaat acgacactaa catgatgaac atgcatgagc 180
tttgaaaagt gctctgtagt cttatgatga tctagaagag cactgtccaa tagaactttc 240
tgtgatgatg aaaagattct acttctgacc tattcaatag ggtaaccact aatcatgcat 300
ggctctcaag cacttgaaat gttgctagtg tgattgggga gctgcgtttt gaatgttaac 360
naatttanat tttaaatcnt taaaaagttt acatqtqqqt taqtqgqncg ccgtacq
```

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<210> 378
<211> 439
<212> DNA
<213> Homo sapiens
<223> Genbank Accession No. H97538
<400> 378
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cttcatctct ttccattttg cggacactcc ccttcttcta ttctccttta ctcaaaacat 120
atggtttaga cccacatcat ggctttcttg tgggaagcct ggatgggact aggaaaacac 180
atgtttccaa catggtgcat atctgtttgt gcagatatca gacaagattt aatcttgtct 240
aacttatgcg tattgttttg atgtttgcct gtggttattc tgggcacagc aatggtggac 300
attattqaaa atqaacttta ttgqcaqatq aaaqataata qaacatqaaq atttatqaac 360
taccataagc tctgcatctc tgggtcttca tttccaaagc agcacttgga aaaccaagcc 420
cagtttcagg caaagagtt
                                                                   439
<210> 379
<211> 440
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H98835
<220>
<221> unsure
<222> (1)..(440)
<223> n = a or c or g or t
<400> 379
caagatcctg cctcccaagc ctataagctt taccaggaga gaggcaggcc ccaccccaag 60
atccactatc cactetttga agaaagatta gagccatgtt ctcagacttt gggctgcatc 120
ctaatccctg cgaagtgcac aatgtgtgat gactccaccc tccacccgat ccagagggtc 180
tggggtgaga cccaaggctg agaggcctcg atggcttcct ggccccatct ccggcagcag 240
ctctatggct gggctctcct gcaggctggg tgcaccccag gccctcagat ggttctaacc 300
agaatcgatg ggcagcagtg acttcgactg tatcatcaat cttggctgcc acaaggttgg 360
gttgtccagg ccctcagctt ganccttqqa ggtgqqccc ccacacaqaq ctttqtctqc 420
ccccagccca ccctcattta
<210> 380
<211> 495
<212> DNA
<213> Homo sapiens
<220×
<223> Genbank Accession No. H99035
<220>
<221> unsure
<222> (1)..(495)
<223> n = a or c or g or t
<400> 380
tgagctttgg acaaatttat tgaaacatac aggcggctgt tagcagagaa atcattccat 60
```

```
gattgatgtg ttacatttgg ccactacctt gaatgtataa tttaaaaatt atatttttca 120
caactaagcc tttgncaaaa aagtcattta gcacatcttt aaagatcaat aagaaatgga 180
tttttggacat taaaaagatc aagtcactga attaaacagt agcaaccccc attaatctag 240
aatcccatag tgctgaaggt agaggtgtct gtgcaaagct agtcatttgt taacagcaat 300
ggctgggctg gtctggtcag gtgagcatgt cccagagaca gcagcaacag agagccgtcc 420
agcaggetgt gaggeaggtg gatggteeta geteatetee teettgggte ttetaceaea 480
tacactgtgg gnttt
                                                                 495
<210> 381
<211> 424
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H99648
<220>
<221> unsure
<222> (1)..(424)
<223> n = a or c or g or t
<400> 381
ggggtatata attttatttt aagtttatat ttcctgcagg atagcaacat acatcttttc 60
ctacccagag gcaaaataca ttttccaaaa acgtggacac tgcccactgc attaagttta 120
aagtgctccc tatatatata gacagtaaaa gtaagcaaag aaacttacaa cacattccaa 180
tctttaatat ctcaaaaatg tttccaaggc aacattatta aaataattat accacagtcc 240
ctaatataac atcaagctcc agtaggaagg tacagagagg gcaggaagtt tccatccagt 300
ctggtttagg tgctcttctt ttcttcaccc agtaaattca cqqtaqcttt cttcqcttct 360
ttagtgatgg catctgcagt ccccttggcc ntgtctttaa gggtccctga ccacactggt 420
ccat
<210> 382
<211> 438
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. H99694
<220>
<221> unsure
<222> (1)..(438)
\langle 223 \rangle n = a or c or g or t
<400> 382
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agatgaggaa acatgtttgc atctcacact agtgcagaga ttctgaaaaa gaccccactt 120
ggaataccaa accacacatt agattgttct gttcccaatt gtgtgccaaa gtgcactctg 180
aactgttttg gtaaagccga ccgtggagtc atatgaggct gaataacttg ggaqaatqta 240
agtotgcaaa ataaacotag gactggattg atcotcaggo cacttggcag gtgaatgtot 300
cgggagtgaa tatgagacaa gcttcctgaa aaggcttata tgacttaaag aactttttgt 360
ttaagtgttt ggtcccaaat aaactattaa gatatataaa gtaattcact gctcaaaaat 420
taccgtcaga taaatatn
                                                                 438
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<210> 383

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<211> 749
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. J00073
<400> 383
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aaatagaggt gagttcatat aacttqattg qccatattat ttcqtggtat qacatatccc 120
acattatage gaattaatat ctaatqqttt ttctqtqaat cctcccaatq tqttatttqc 180
tecettgett ggaaetteag agtteaetgg aagtttttgt tttettetge agattattge 240
tccccctgag cgtaaatact ctgtctggat tgggggctcc atcttggcct ctctgtccac 300
cttccagcaa atgtggatta gcaagcaaga gtacgatgag gcaggcccat ccattgtcca 360
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tatgettett ggagtetece aaaccacett eceteatett teatcaatea ttgtacagtt 480
tgtttacaca cgtgcaattt gtttgtgctt ctaatattta ttgctttata aataaaccag 540
actaggactt gcaacctata aaagcetete gtttgttttt ggggtaggeg tggggtgggg 600
caggtgtttg ctttgacacc ctgagcattg tcaaagttca gtagcacaay gttcatccag 660
atgaattaat atgacagtta gcrgggagtt ataatgctaa ctttgattca tatttggaca 720
gaatcatgaa tatattcata tccgaagcg
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<210> 384
<211> 1056
<212> DNA
<213> Homo sapiens
<220>
<223> Genbank Accession No. J00123
<400> 384
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gcttcctcaa gatggcacca gcaccctcag agaaaatagc aaaccggaag aaagccattt 180
gctagccaaa aggtatgggg gcttcatgaa aaggtatgga ggcttcatga agaaaatgga 240
tgagetttat cccatggage cagaagaaga ggccaatgga agtgagatcc tcgccaagcg 300
gtatgggggc ttcatgaaga aggatgcaga ggaggacgac tcgctggcca attcctcaga 360
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cagtgataat gaggaagaag tgagcaagag atatgggggc ttcatgagag gcttaaagag 480
aagcccccaa ctggaagatg aagccaaaga gctgcagaag cgatatgggg gcttcatgag 540
aagagtaggt cgcccagagt ggtggatgga ctaccagaaa cggtatggag gtttcctgaa 600
gcgctttgcc gaggctctgc cctccgacga agaaggcgaa agttactcca aagaagttcc 660
tgaaatggaa aaaagatacg gaggatttat gagattttaa tatcttttcc cactagtggc 720
ccccaggccc cagcaagcct ccctccatcc tccagtggga aactgttgat ggtgttttat 780
tgtcatgtgt tgcttgcctt gtatagttga cttcattgtc tggataacta tacaacctga 840
aaactgtcat ttcaggttct gtgctctttt tggagtcttt aagctcagta ttagtctatt 900
gcagctatct cgtttttcat gctaaaaata gttttttgtt atcttgtctc ttattttttg 960
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<210> 385
<211> 1089
<212> DNA
<213> Homo sapiens
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141

<220>

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<220>
<221> unsure
<222> (1)..(1089)
<223> n = a or c or g or t
<400> 385
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teccagatgg gteetgteec aggtgeacet geaggagteg ggeecaggae tggggaagee 120
tccagagctc aaaaccccac ttqqtqacac aactcacaca tqcccacqqt qcccaqaqcc 180
caaatcttgt gacacacctc ccccgtgccc acggtgccca gagcccaaat cttgtgacac 240
acctcccca tgcccacggt gcccagagcc caaatcttgt gacacacctc ccccgtgccc 300
nnngtgccca gcacctgaac tettgggagg accgtcagtc ttcctcttcc ccccaaaacc 360
caaggatacc cttatgattt cccggacccc tgaqqtcacg tgcgtqqtqq tggacqtqag 420
ccacgaagac ccnnnngtcc agttcaagtg gtacgtggac ggcgtggagg tgcataatgc 480
caagacaaag ctgcgggagg agcagtacaa cagcacgttc cgtgtggtca gcgtcctcac 540
cgtcctgcac caggactggc tgaacggcaa ggagtacaag tgcaaggtct ccaacaaagc 600
cctcccagcc cccatcgaga aaaccatctc caaagccaaa ggacagcccn nnnnnnnnn 660
nnnnnnnnn nnnnnnnnn nnnnngagga gatgaccaag aaccaagtca gcctgacctg 720
cctggtcaaa ggcttctacc ccagcgacat cgccgtggag tgggagagca atgggcagcc 780
ggagaacaac tacaacacca cgcctcccat gctggactcc gacggctcct tcttcctcta 840
cagcaagctc accgtggaca agagcaggtg gcagcagggg aacatcttct catgctccgt 900
gatgcatgag gctctgcaca accgctacac gcagaaqagc ctctccctqt ctccqqqtaa 960
atgagtgcca tggccggcaa gccccgctc cccgggctct cggggtcgcg cgaggatgct 1020
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<211> 2133
<212> DNA
<213> Homo sapiens
<220>
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